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DESCRIPTION

POLYHYDROXYALKANOATE HAVING ESTER GROUP, CARBOXYL GROUP, AND SULFONIC GROUP, AND METHOD OF PRODUCING THE SAME

TECHNICAL FIELD

The present invention relates to a novel polyhydroxyalkanoate and a method of producing the same.

BACKGROUND ART

Biodegradable polymer materials have been finding a wide variety of applications including medical materials, drug delivery systems, and environmentally compatible materials. In recent years, in addition to those applications, the biodegradable polymer materials have been requested to provide new functions, and hence various studies have been made. In particular, the introduction of a chemically modifiable functional group into a molecule of a polyhydroxyalkanoate typified by polylactic acid has been examined. There has been reported a compound into which a carboxyl group or a vinyl group is introduced. For example, polymalic acid has been known as a polyhydroxyalkanoate having a carboxyl group at a side chain thereof. An α -type

represented by the chemical formula (39) and a β -type represented by the chemical formula (40) have been known as polymers of polymalic acid depending on the form of a polymer.

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Of those, a polymer obtained by ring-opening polymerization of a benzyl ester of β -malolactone represented by the chemical formula (41) is disclosed in US 4,265,247 (Patent Document 1) as β -type polymalic acid or a copolymer thereof.

(R₄₁: benzyl group.)

In addition, a polymer obtained by copolymerization of a six-membered ring diester monomer represented by the chemical formula (42) and a glicolide or lactide as a cyclic diester or a lactone as an intramolecular ring closure reaction ester of ω -hydroxycarboxylic acid is disclosed in JP-A 02-3415 (Patent Document 2) as a copolymer containing any one of other

hydroxyalkanoic acids typified by α -type polymalic acid-glycolic acid copolymer and glycolic acid.

(R₄₂ represents a lower alkyl group such as a methyl group, an ethyl group, an n-propyl group, an isopropyl group, or a t-butyl group, or a benzyl group.)

Macromolecules 2000, 33 (13), 4619-4627 (Non-Patent Document 1) discloses that 7-oxo-4oxepanonecarboxylate is subjected to ring-opening 10 polymerization to produce a polymer having an ester group at a side chain thereof, and the polymer is further subjected to hydrogenolysis to produce a polymer having a carboxylic acid at a side chain thereof as a polyhydroxyalkanoate having a carboxyl 15 group at a side chain thereof. Biomacromolecules 2000, 1, 275 (Non-Patent Document 2) discloses a polymer in which a benzyloxycarbonyl group is introduced into a methylene group at lpha -position of a carbonyl group in the main-chain of poly(ε-20 caprolactone), the polymer being obtained by: allowing lithium diisopropylamide to react with poly(ε -caprolactone); and allowing the resultant to react with benzyl chloroformate. Macromolecular

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Bioscience 2004, 4, 232 (Non-Patent Document 3) discloses a polymer in which a (benzyloxycarbonyl)methyl group is introduced into a methylene group at α —position of a carbonyl group in the main chain of polylactic acid, the polymer being obtained by: allowing lithium diisopropylamide to react with polylactic acid; and allowing the resultant to react with benzyl bromoacetate.

Polymeric Materials Science & Engineering 2002, 87, 254 (Non-Patent Document 4) discloses, as a polyhydroxyalkanoate having a vinyl group at a side chain thereof, a polymer obtained by ring-opening polymerization of α-allyl(δ-valerolactone). Similarly, Polymer Preprints 2002, 43 (2), 727 (Non-Patent Document 5) discloses, as a polyhydroxyalkanoate having a vinyl group at a side chain thereof, a polymer obtained by ring-opening polymerization of 3,6-diallyl-1,4-dioxane-2,5-dione as a six-membered ring diester monomer.

There has been reported a polymer having a new function into which a structure providing functional properties for a polyhydroxyalkanoate into which a chemically modifiable functional group is introduced as described above is introduced. International Journal of Biological Macromolecules 25 (1999) 265 (Non-Patent Document 6) discloses the following. A copolymer of α -type malic acid and glycolic acid is

obtained by ring-opening polymerization of a cyclic dimer of α -type malic acid and glycolic acid, and the resultant polymer is deprotected to obtain a polyester having a carboxyl group at a side chain thereof. Tripeptide is chemically modified to the carboxyl group at the side chain, and the resultant polymer is evaluated for cell adhesion. At this time, a good result is obtained.

10 DISCLOSURE OF THE INVENTION

It may be possible that new functional properties can be provided by introducing a unit having a carboxyl group that is a reactive functional group, in a molecule as described above; and chemically modifying the reactive functional group. 15 However, there have been a small number of reports concerning it. In view of the above, the present invention provides: a novel polyhydroxyalkanoate having a reactive functional group in a molecule and 20 a method of producing the same; and a novel polyhydroxyalkanoate having a new function obtained by chemically modifying the polyhydroxyalkanoate having a reactive functional group and a method of producing the same.

The inventors of the present invention have made extensive studies with a view to developing a novel polyhydroxyalkanoate having a reactive

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functional group in a molecule and a novel polyhydroxyalkanoate having a new function obtained by chemically modifying the polyhydroxyalkanoate having a reactive functional group, thereby completing the invention described below.

The polyhydroxyalkanoate according to the present invention includes the following.

(1) A polyhydroxyalkanoate, comprising one or more units each represented by the chemical formula
10 (1).

$$\begin{array}{c}
R \\
N-H \\
= O \\
(CH_2)m \\
\hline
\begin{pmatrix}
1 \\
O \\
Z_{1b}
\end{pmatrix} = Z_{1a} - O \\
\end{array}$$
(1)

(In the formula, R represents $-A_1-SO_2R_1$. R_1 represents OH, a halogen atom, ONa, OK, or OR_{1a} . R_{1a} and A_1 each independently represent a group having a substituted or unsubstituted aliphatic hydrocarbon structure, a substituted or unsubstituted aromatic ring structure, or a substituted or unsubstituted heterocyclic structure. Z_{1a} represents a linear alkylene chain having 1 to 4 carbon atoms. The linear alkylene chain has at least one linear or branched alkyl group, or at least one alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal

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thereof. Z_{1b} represents a hydrogen atom, a linear or branched alkyl group, aryl group, or aralkyl group which may be substituted by an aryl group. In addition, m represents an integer selected from 0 to 8. When multiple units exist, R, R_1 , R_{1a} , A_1 , Z_{1a} , Z_{1b} , and m each independently have the above meaning for each unit.)

(2) A polyhydroxyalkanoate, comprising one or more units each represented by the chemical formula (5).

(In the formula, R_5 represents hydrogen, a group for forming a salt, or R_{5a} . R_{5a} represents a linear or branched alkyl or aralkyl group having 1 to 12 carbon atoms, or a group having a saccharide. m represents an integer selected from 0 to 8. Z_{5a} represents a linear alkylene chain having 1 to 4 carbon atoms. The linear alkylene chain has at least one linear or branched alkyl group, or at least one alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof. Z_{5b} represents a hydrogen atom, or a linear or branched alkyl group, aryl group, or aralkyl group which may be substituted

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by an aryl group. When multiple units exist, R_5 , R_{5a} , Z_{5a} , Z_{5b} , and m each independently have the above meaning for each unit.)

On the other hand, the method of producing

5 polyhydroxyalkanoate according to the present
invention includes the following.

(A) A method of producing a polyhydroxyalkanoate containing a unit represented by the chemical formula (1), characterized by comprising the step of subjecting a polyhydroxyalkanoate containing a unit represented by the chemical formula (29) and at least one kind of amine compound represented by the chemical formula (30) to a condensation reaction.

$$\begin{array}{c} COOR_{29} \\ (CH_2)m \\ \hline \\ O \\ Z_{29b} \end{array} (29)$$

(In the formula, R_{29} represents hydrogen or a group for forming a salt. m represents an integer selected from 0 to 8. Z_{29a} represents a linear alkylene chain having 1 to 4 carbon atoms. The linear alkylene chain has at least one linear or branched alkyl group, or at least one alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof. Z_{5b} represents a hydrogen atom, or a linear

or branched alkyl group, aryl group, or aralkyl group which may be substituted by an aryl group. When multiple units exist, R_{29} , Z_{29a} , Z_{29b} , and m each independently have the above meaning for each unit.) $H_2N - A_3 - SO_2R_{30}$ (30)

(In the formula, R₃₀ represents OH, a halogen atom, ONa, OK, or OR_{30a}. R_{30a} and A₃ each independently represent a group having a substituted or unsubstituted aliphatic hydrocarbon structure, a substituted or unsubstituted aromatic ring structure, or a substituted or unsubstituted heterocyclic structure. When multiple units exist, R₃₀, R_{30a}, and A₃, and m each independently have the above meaning for each unit.)

$$\begin{array}{c}
R\\N-H\\ = O\\ (CH_2)m\\ \hline
\begin{pmatrix}
II & -Z_{1a} & -O
\end{pmatrix}
\end{array}$$

$$\begin{array}{c}
C & -Z_{1a} & -O
\end{array}$$
(1)

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(In the formula, R represents $-A_1-SO_2R_1$. R_1 represents OH, a halogen atom, ONa, OK, or OR_{1a} . R_{1a} and A_1 each independently represent a group having a substituted or unsubstituted aliphatic hydrocarbon structure, a substituted or unsubstituted aromatic ring structure, or a substituted or unsubstituted heterocyclic structure. Z_{1a} represents a linear alkylene chain having 1 to 4 carbon atoms. The linear alkylene

chain has at least one linear or branched alkyl group, or at least one alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof. Z_{1b} represents a hydrogen atom, or a linear or branched alkyl group, aryl group, or aralkyl group which may be substituted by an aryl group. In addition, m represents an integer selected from 0 to 8. When multiple units exist, R, R₁, R_{1a}, A₁, Z_{1a}, Z_{1b}, and m each independently have the above meaning for each unit.)

(B) A method of producing a polyhydroxyalkanoate containing a unit represented by the chemical formula (32), characterized by

15 comprising the step of hydrolyzing a polyhydroxyalkanoate containing a unit represented by the chemical formula (31) in the presence of an acid or an alkali, or the step of subjecting the polyhydroxyalkanoate to hydrogenolysis including

20 catalytic reduction.

$$COOR_{31}$$

$$(CH_2)m$$

$$Z_{31a}-O$$

$$Z_{31b}$$

$$(31)$$

(In the formula, R_{31} represents a linear or branched alkyl or aralkyl group having 1 to 12 carbon atoms. Z_{31a} represents a linear alkylene chain having 1 to 4

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carbon atoms. The linear alkylene chain has at least one linear or branched alkyl group, or at least one alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof. Z_{31b} represents a hydrogen atom, or a linear or branched alkyl group, aryl group, or aralkyl group which may be substituted by an aryl group. In addition, m represents an integer selected from 0 to 8. When multiple units exist, R₃₁, Z_{31a}, Z_{31b}, and m each independently have the above meaning for each unit.)

(In the formula, R_{32} represents hydrogen or a group for forming a salt. Z_{32a} represents a linear alkylene chain having 1 to 4 carbon atoms. The linear alkylene chain has at least one linear or branched alkyl group, or at least one alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof. Z_{32b} represents a hydrogen atom, or a linear or branched alkyl group, aryl group, or aralkyl group which may be substituted by an aryl group. m represents an integer selected from 0 to 8. When multiple units exist, R_{32} , Z_{32a} , Z_{32b} , and m each

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independently have the above meaning for each unit.)

(C) A method of producing a polyhydroxyalkanoate containing a unit represented by the chemical formula (35), characterized by including the steps of: allowing a polyhydroxyalkanoate containing a unit represented by the chemical formula (33) to react with a base; and allowing the compound obtained in the foregoing step to react with a compound represented by the chemical formula (34).

$$\begin{array}{c|c}
H \\
\hline
O \\
Z_{33b}
\end{array}$$
(33)

(In the formula, Z_{33a} represents a linear alkylene chain having 1 to 4 carbon atoms. The linear alkylene chain has at least one linear or branched alkyl group, or at least one alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof. Z_{33b} represents a hydrogen atom, or a linear or branched alkyl group, aryl group, or aralkyl group which may be substituted by an aryl group. When multiple units exist, Z_{33a} and Z_{33b} each independently have the above meaning for each unit.) X(CH₂)mCOOR₃₄

(In the formula, m represents an integer selected from 0 to 8. X represents a halogen atom. R_{34} represents a linear or branched alkyl or aralkyl

(34)

group having 1 to 12 carbon atoms.)

$$\begin{array}{c}
COOR_{35} \\
(CH_2)m \\
\hline
COOR_{35} \\
\hline
Z_{35a} - O
\end{array}$$
(35)

(In the formula, R₃₅ represents a linear or branched alkyl or aralkyl group having 1 to 12 carbon atoms. Z_{35a} represents a linear alkylene chain having 1 to 4 5 carbon atoms. The linear alkylene chain has at least one linear or branched alkyl group, or at least one alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof. Z35b 10 represents a hydrogen atom, or a linear or branched alkyl group, aryl group, or aralkyl group which may be substituted by an aryl group. In addition, m represents an integer selected from 0 to 8. When multiple units exist, R₃₅, Z_{35a}, Z_{35b}, and m each 15 independently have the above meaning for each unit.)

(D) A method of producing a polyhydroxyalkanoate containing a unit represented by the chemical formula (38), characterized by including the steps of: allowing a polyhydroxyalkanoate containing a unit represented by the chemical formula (36) to react with a base; and allowing the compound obtained in the foregoing step to react with a compound represented by the chemical formula (37).

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(In the formula, Z_{36a} represents a linear alkylene chain having 1 to 4 carbon atoms. The linear alkylene chain has at least one linear or branched alkyl group, or at least one alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof. Z_{36b} represents a hydrogen atom, or a linear or branched alkyl group, aryl group, or aralkyl group which may be substituted by an aryl group. When multiple units exist, Z_{36a} and Z_{36b} each independently have the above meaning for each unit.)

(In the formula, R₃₇ represents -A₃₇-SO₂R_{37a}. R_{37a}

15 represents OH, a halogen atom, ONa, OK, or OR_{37b}. R_{37b}

and A₃₇ are each independently selected from groups
each having a substituted or unsubstituted aliphatic
hydrocarbon structure, a substituted or unsubstituted
aromatic ring structure, or a substituted or

20 unsubstituted heterocyclic structure. When multiple
units exist, R₃₇, R_{37a}, R_{37b}, and A₃₇ each independently
have the above meaning for each unit.)

(In the formula, R₃₈ represents -A₃₈-SO₂R_{38a}. R_{38a} represents OH, a halogen atom, ONa, OK, or OR38b. and A38 each independently represent a group having a substituted or unsubstituted aliphatic hydrocarbon 5 structure, a substituted or unsubstituted aromatic ring structure, or a substituted or unsubstituted heterocyclic structure. Z38a represents a linear alkylene chain having 1 to 4 carbon atoms. linear alkylene chain has at least one linear or 10 branched alkyl group, or at least one alkyl group containing a residue having any one of a phenylstructure, a thienyl structure, and a cyclohexyl structure at a terminal thereof. Z_{38b} represents a hydrogen atom, or a linear or branched alkyl group, 15 aryl group, or aralkyl group which may be substituted by an aryl group. When multiple units exist, R_{38} , R_{38a} , R_{38b}, A₃₈, Z_{38a}, and Z_{38b} each independently have the above meaning for each unit.)

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BEST MODE FOR CARRYING OUT THE INVENTION

Hereinafter, the contents of the present invention will be described. A polyhydroxyalkanoate

containing a unit represented by the chemical formula (1) as a target in the present invention can be produced by a reaction between a polyhydroxyalkanoate containing a unit represented by the chemical formula (29) to be used as a starting material and at least one kind of aminosulfonic acid compound represented by the chemical formula (30).

$$\begin{array}{c|c}
COOR_{29} \\
(CH_2)m \\
\hline
COOR_{29a} - O
\end{array}$$

$$\begin{array}{c|c}
Z_{29a} - O
\end{array}$$
(29)

(In the formula, R_{29} represents hydrogen or a group 10 for forming a salt. m represents an integer selected from 0 to 8. Z_{29a} represents a linear alkylene chain having 1 to 4 carbon atoms. The linear alkylene chain has at least one linear or branched alkyl group, or at least one alkyl group containing a residue having any one of a phenyl structure, a thienyl 15 structure, and a cyclohexyl structure at a terminal thereof. Z_{29b} represents a hydrogen atom, or a linear or branched alkyl group, aryl group, or aralkyl group which may be substituted by an aryl group. When multiple units exist, R_{29} , Z_{29a} , Z_{29b} , and m each 20 independently have the above meaning for each unit.)

 H_2N — A_3 — SO_2R_{30} (30) (In the formula, R_{30} represents OH, a halogen atom,

ONa, OK, or OR_{30a} . R_{30a} and A_3 are each independently

selected from groups each having a substituted or unsubstituted aliphatic hydrocarbon structure, a substituted or unsubstituted aromatic ring structure, or a substituted or unsubstituted heterocyclic structure (R_{30a} represents a monovalent group having a structure selected from them, and A_3 represents a divalent group having a structure selected from them). When multiple units exist, R_{30} , R_{30a} , and A_3 each independently have the above meaning for each unit.)

More specifically, in the compound represented by the chemical formula (29) to be used in the present invention, Z₂₉ represents a linear alkylene chain having 1 to 4 carbon atoms. The linear alkylene chain has at least one linear or branched alkyl group, or at least one alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof. The linear alkylene chain structure represented by Z₂₉ is preferably selected from the following (A) to (D).

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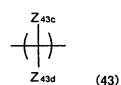
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(A) When the linear alkylene chain has 1 carbon atom, in the linear alkylene chain structure represented by the chemical formula (43), one of Z_{43c} and Z_{43d} represents a linear or branched alkyl group, or an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof.

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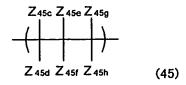
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(B) When the linear alkylene chain has 2 carbon atoms, in the linear alkylene chain structure represented by the chemical formula (44), one of Z_{44c} , Z_{44d} , Z_{44e} , and Z_{44f} represents a linear or branched alkyl group, or an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof.

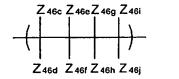


(C) When the linear alkylene chain has 3 carbon atoms, in the linear alkylene chain structure represented by the chemical formula (45), one of Z_{45c} , Z_{45d} , Z_{45e} , Z_{45f} , Z_{45g} , and Z_{45h} represents a linear or branched alkyl group, or an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof.



(D) When the linear alkylene chain has 4 carbon atoms, in the linear alkylene chain structure

represented by the chemical formula (46), one of Z_{46c} , Z_{46d} , Z_{46e} , Z_{46f} , Z_{46g} , Z_{46h} , Z_{46i} , and Z_{46h} represents a linear or branched alkyl group, or an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof.



When a substituent selected from Z_{43c}, Z_{43d}, Z_{44c}, Z_{44d}, Z_{44e}, Z_{44f}, Z_{45c}, Z_{45d}, Z_{45e}, Z_{45f}, Z_{45g}, Z_{45h}, Z_{46c}, Z_{46d}, Z_{46e}, Z_{46f}, Z_{46g}, Z_{46h}, Z_{46i}, and Z_{46j} described in the chemical formulae (43), (44), (45), and (46) represents a linear or branched alkyl group, or an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof, the substituent is more preferably selected from substituents represented by the chemical formulae (14), (15), (16), and (17).

—(CH₂)k₁₄-CH₃ (14)

(46)

20 (In the formula, k_{14} represents an integer selected from 0 to 8. When multiple units exist, k_{14} 's each independently have the above meaning for each unit.)

$$--(CH2)k15---(CH3)CH3 (15)$$

(In the formula, k_{15} represents an integer selected

from 0 to 7. When multiple units exist, k_{15} 's each independently have the above meaning for each unit.) $-(CH_2)k_{\overline{16}}-R_{16}$ (16)

(In the formula, k_{16} represents an integer selected from 1 to 8. R_{16} represents a substituent containing a residue having any one of a phenyl structure and a thienyl structure. When multiple units exist, k_{15} and R_{16} each independently have the above meaning for each unit.)

$$-(CH_2)k_{17}$$

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(In the formula, R_{17} represents a substituent to a cyclohexyl group selected from an H atom, a CN group, an NO_2 group, a halogen atom, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a CF_3 group, a C_2F_5 group, and a C_3F_7 group. k_{17} represents an integer selected from 0 to 8. When multiple units exist, k_{17} and R_{17} each independently have the above meaning for each unit.)

In addition, R_{16} in the chemical formula (16), that is, a residue having any one of a phenyl structure and a thienyl structure is selected from the group of residues represented by the chemical formulae (18), (19), (20), (21), (22), (23), (24), (25), (26), (27), and (28).

Here, the chemical formula (18) represents a group of unsubstituted or substituted phenyl groups.

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(In the formula, R_{18} represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $CH=CH_2$ group, $COOR_{18a}$ (R_{18a} represents an H atom, an Na atom, or a K atom.), a CF_3 group, a C_2F_5 group, and a C_3F_7 group. When multiple units exist, R_{18} 's may be different for each unit.)

The chemical formula (19) represents a group of unsubstituted or substituted phenoxy groups.

(In the formula, R_{19} represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a SCH_3 group, a CF_3 group, a C_2F_5 group, and a C_3F_7 group. When multiple units exist, R_{19} 's may be different for each unit.)

The chemical formula (20) represents a group of unsubstituted or substituted benzoyl groups.

(In the formula, R_{20} represents a substituent to an

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aromatic ring selected from an H atom, a halogen atom, a CN group, an NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a CF_3 group, a C_2F_5 group, and a C_3F_7 group. When multiple units exist, R_{20} 's may be different for each unit.)

The chemical formula (21) represents a group of unsubstituted or substituted phenylsulfanyl groups.

(In the formula, R_{21} represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO_2 group, $COOR_{21a}$, SO_2R_{21b} (R_{21a} represents H, Na, K, CH₃, or C_2H_5 , and R_{21b} represents OH, ONa, OK, a halogen atom, OCH₃, or OC_2H_5 .), a CH₃ group, a C_2H_5 group, a C_3H_7 group, a $(CH_3)_2$ -CH group, and a $(CH_3)_3$ -C group. When multiple units exist, R_{21} 's may be different for each unit.)

The chemical formula (22) represents a group of unsubstituted or substituted (phenylmethyl)sulfanyl groups.

(In the formula, R_{22} represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO_2 group, $COOR_{22a}$, SO_2R_{22b} (R_{22a}

represents H, Na, K, CH₃, or C_2H_5 , and R_{22b} represents OH, ONa, OK, a halogen atom, OCH₃, or OC₂H₅.), a CH₃ group, a C_2H_5 group, a C_3H_7 group, a (CH₃)₂-CH group, and a (CH₃)₃-C group. When multiple units exist, R_{22} 's may be different for each unit.)

The chemical formula (23) represents a 2-thienyl group.

The chemical formula (24) represents a 2-

10 thienylsulfanyl group.

The chemical formula (25) represents a 2-thienylcarbonyl group.

The chemical formula (26) represents a group of unsubstituted or substituted phenylsulfinyl groups.

(In the formula, R₂₆ represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO₂ group, COOR_{26a}, SO₂R_{26b} (R_{26a} represents H, Na, K, CH₃, or C₂H₅, and R_{26b} represents OH, ONa, OK, a halogen atom, OCH₃, or OC₂H₅.), a CH₃

group, a C_2H_5 group, a C_3H_7 group, a $(CH_3)_2$ -CH group, and a $(CH_3)_3$ -C group. When multiple units exist, R_{26} 's may be different for each unit.)

The chemical formula (27) represents a group of unsubstituted or substituted phenylsulfonyl groups.

(In the formula, R₂₇ represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO₂ group, COOR_{27a}, SO₂R_{27b} (R_{27a}

10 represents H, Na, K, CH₃, or C₂H₅, and R_{27b} represents OH, ONa, OK, a halogen atom, OCH₃, or OC₂H₅.), a CH₃ group, a C₂H₅ group, a C₃H₇ group, a (CH₃)₂-CH group, and a (CH₃)₃-C group. When multiple units exist, R₂₇'s may be different for each unit.)

The chemical formula (28) represents a (phenylmethyl)oxy group.

On the other hand, in the compound represented by the chemical formula (30) to be used in the 20 present invention, R₃₀ represents OH, a halogen atom, ONa, OK, or OR₃₀. R₃₀ represents a linear or branched alkyl group having 1 to 8 carbon atoms, or a substituted or unsubstituted phenyl group.

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A₃ represents a liner or branched and substituted or unsubstituted alkylene group having 1 to 8 carbon atoms, a substituted or unsubstituted phenylene group, a substituted or unsubstituted naphthalene group, or a substituted or unsubstituted heterocyclic structure containing one or more of N, S, and O. When A₃ represents a ring structure, an unsubstituted ring may be further condensed. In addition, when multiple units exist, R₃₀, R_{30a}, and A₃ each independently have the above meaning for each unit.

When A_3 represents a linear and substituted or unsubstituted alkylene group, an aminosulfonic acid compound represented by the following chemical

15 formula (47) is exemplified.

$$H_2N - A_4 - SO_2R_{47}$$
 (47)

(In the formula, R_{47} represents OH, a halogen atom, ONa, OK, or OR_{47a} . R_{47a} represents a linear or branched alkyl group having 1 to 8 carbon atoms, or a substituted or unsubstituted phenyl group. A_4 represents a liner or branched and substituted or unsubstituted alkylene group having 1 to 8 carbon atoms, which may be substituted by an alkyl group having 1 to 20 carbon atoms, an alkoxy group having 1 to 20 carbon atoms, or the like as a substituent.)

Examples of the compound represented by the chemical formula (47) include 2-aminoethanesulfonic

acid (taurine), 3-aminopropanesulfonic acid, 4-aminobutanesulfonic acid, 2-amino-2-methylpropanesulfonic acid, and alkali metal salts and esterified products thereof.

When A₃ represents a substituted or unsubstituted phenylene group, an aminosulfonic acid compound represented by the following chemical formula (48) is exemplified.

(In the formula, R_{3a} , R_{3b} , R_{3c} , R_{3d} , and R_{3e} each 10 independently represent SO_2R_{3f} (R_{3f} represents OH, a halogen atom, ONa, OK, or OR_{3f1}. (R_{3f1} represents a linear or branched alkyl group having 1 to 8 carbon atoms, or a substituted or unsubstituted phenyl group.)), a hydrogen atom, a halogen atom, an alkyl 15 group having 1 to 20 carbon atoms, an alkoxy group having 1 to 20 carbon atoms, an OH group, an NH_2 group, an NO2 group, COOR3q (R3q represents an H atom, an Na atom, or a K atom.), an acetamide group, an OPh group, an NHPh group, a CF3 group, a C2F5 group, or a 20 C₃F₇ group (Ph represents a phenyl group.), and at least one of these groups represents SO₂R_{3f}.)

A polyhydroxyalkanoate having one or more units each represented by the chemical formula (3) can be

obtained by using a compound represented by the chemical formula (48).

$$\begin{array}{c|c}
R_{3b} & R_{3c} \\
R_{3a} & R_{3e} \\
\hline
R_{3e} & R_{3e}
\end{array}$$

$$\begin{array}{c|c}
CCH_2 & CCH_2 & CCH_2 \\
\hline
CCH_2 & CCH_2 & CCH_2
\end{array}$$

$$\begin{array}{c|c}
CCH_2 & CCH_2 & CCH_2 & CCH_2
\end{array}$$

$$\begin{array}{c|c}
CCH_2 & CCH_2 & CCH_2 & CCH_2
\end{array}$$

$$\begin{array}{c|c}
CCH_2 & CCH_2 & CCH_2
\end{array}$$

$$\begin{array}{c|c}
CCH_2 & CCH_2 & CCH_2
\end{array}$$

$$\begin{array}{c|c}
CCH_2 & CCH_2
\end{array}$$

(In the formula, R_{3a} , R_{3b} , R_{3c} , R_{3d} , and R_{3e} each independently represent SO_2R_{3f} (R_{3f} represents OH, a 5 halogen atom, ONa, OK, or OR3f1. (R3f1 represents a linear or branched alkyl group having 1 to 8 carbon atoms, or a substituted or unsubstituted phenyl group.)), a hydrogen atom, a halogen atom, an alkyl group having 1 to 20 carbon atoms, an alkoxy group 10 having 1 to 20 carbon atoms, an OH group, an NH_2 group, an NO_2 group, $COOR_{3g}$ (R_{3g} represents an H atom, an Na atom, or a K atom.), an acetamide group, an OPh group, an NHPh group, a CF3 group, a C2F5 group, or a C₃F₇ group (Ph represents a phenyl group.), and at 15 least one of these groups represents SO₂R_{3f}. Z_{3a} represents a linear alkylene chain having 1 to 4. carbon atoms. The linear alkylene chain has at least one linear or branched alkyl group, or at least one alkyl group containing a residue having any one of a 20

phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof. Z_{3b} represents a hydrogen atom, or a linear or branched alkyl group, aryl group, or aralkyl group which may be substituted by an aryl group. In addition, m represents an integer selected from 0 to 8. When multiple units exist, R_{3a} , R_{3b} , R_{3c} , R_{3d} , R_{3e} , R_{3f} , R_{3f1} , R_{3g} , Z_{1a} , Z_{1b} , and m each independently have the above meaning for each unit.)

Examples of the compound represented by the 10 chemical formula (48) include p-aminobenzenesulfonic acid (sulfanilic acid), m-aminobenzenesulfonic acid, o-aminobenzenesulfonic acid, m-toluidine-4-sulfonic acid, sodium o-toluidine-4-sulfonate, p-toluidine-2sulfonic acid, 4-methoxyaniline-2-sulfonic acid, o-15 anisidine-5-sulfonic acid, p-anisidine-3-sulfonic acid, 3-nitroaniline-4-sulfonic acid, sodium 2nitroaniline-4-sulfonate, sodium 4-nitroaniline-2sulfonate, 1,5-dinitroaniline-4-sulfonic acid, 2aminophenol-4-hydroxy-5-nitrobenzenesulfonic acid, 20 sodium 2,4-dimethylaniline-5-sulfonate, 2,4dimethylaniline-6-sulfonic acid, 3,4-dimethylaniline-5-sulfonic acid, 4-isopropylaniline-6-sulfonic acid, 4-trifluoromethylaniline-6-sulfonic acid, 3-carboxy-4-hydroxyaniline-5-sulfonic acid, 4-carboxyaniline-6-25 sulfonic acid, and alkali metal salts and esterified products thereof.

When A_3 represents a substituted or unsubstituted naphthalene group, an aminosulfonic acid compound represented by the following chemical formula (49A) or (49B) is exemplified.

$$R_{4g}$$
 R_{4g}
 R_{4d}
 R_{4c}
 R_{4b}
 R_{4b}
 R_{4b}
 R_{4b}
 R_{4b}
 R_{4b}

(In the formula, R_{4a}, R_{4b}, R_{4c}, R_{4d}, R_{4e}, R_{4f}, and R_{4g} each independently represent SO₂R_{4o} (R_{4o} represents OH, a halogen atom, ONa, OK, or OR_{4o1}. (R_{4o1} represents a linear or branched alkyl group having 1 to 8 carbon atoms, or a substituted or unsubstituted phenyl group.)), a hydrogen atom, a halogen atom, an alkyl group having 1 to 20 carbon atoms, an alkoxy group having 1 to 20 carbon atoms, an OH group, an NH₂ group, an NO₂ group, COOR_{4p} (R_{4p} represents an H atom, an Na atom, or a K atom.), an acetamide group, an OPh group, an NHPh group, a CF₃ group, a C₂F₅ group, or a C₃F₇ group (Ph represents a phenyl group.), and at least one of these groups represents SO₂R_{4o}.)

20 (In the formula, R_{4n} , R_{4i} , R_{4j} , R_{4k} , R_{4l} , R_{4m} , and R_{4n}

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each independently represent SO_2R_{4o} (R_{4o} represents OH, a halogen atom, ONa, OK, or OR_{4o1} . (R_{4o1} represents a linear or branched alkyl group having 1 to 8 carbon atoms, or a substituted or unsubstituted phenyl group.)), a hydrogen atom, a halogen atom, an alkyl group having 1 to 20 carbon atoms, an alkoxy group having 1 to 20 carbon atoms, an OH group, an NH_2 group, an NO_2 group, $COOR_{4p}$ (R_{4p} represents an H atom, an Na atom, or a K atom.), an acetamide group, an OPh group, an NHPh group, a CF_3 group, a C_2F_5 group, or a C_3F_7 group (Ph represents a phenyl group.), and at least one of these groups represents SO_2R_{4o} .)

A polyhydroxyalkanoate having one or more units each represented by the chemical formula (4A) or (4B) can be obtained by using a compound represented by the chemical formula (49A) or (49B).

$$R_{4g}$$
 R_{4g}
 R_{4g}
 R_{4b}
 R

(In the formula, R_{4a} , R_{4b} , R_{4c} , R_{4d} , R_{4e} , R_{4f} , and R_{4g} each independently represent SO_2R_{4o} (R_{4o} represents OH,

a halogen atom, ONa, OK, or OR401. (R401 represents a linear or branched alkyl group having 1 to 8 carbon atoms, or a substituted or unsubstituted phenyl group.)), a hydrogen atom, a halogen atom, an alkyl group having 1 to 20 carbon atoms, an alkoxy group having 1 to 20 carbon atoms, an OH group, an NH2 group, an NO₂ group, COOR_{4p} (R_{4p} represents an H atom, an Na atom, or a K atom.), an acetamide group, an OPh group, an NHPh group, a CF3 group, a C2F5 group, or a C₃F₇ group (Ph represents a phenyl group.), and at least one of these groups represents SO_2R_{40} . Z_{1a} represents a linear alkylene chain having 1 to 4 carbon atoms. The linear alkylene chain has at least one linear or branched alkyl group, or at least one alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof. Z_{1b} represents a hydrogen atom, or a linear or branched alkyl group, aryl group, or aralkyl group which may be substituted by an aryl group. In addition, m represents an integer selected from 0 to 8. When multiple units exist, R_{4a}, R_{4b}, R_{4c}, R_{4d}, R_{4e}, R_{4f}, R_{4g}, R_{40} , R_{401} , R_{4p} , m, Z_{1a} , Z_{1b} , and n each independently have the above meaning for each unit.)

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$$\begin{array}{c|c}
R_{4h} & R_{4i} \\
R_{4m} & R_{4h} \\
R_{4h} & R_{4h}
\end{array}$$

$$\begin{array}{c|c}
CH_2 \\
C \\
Z_{1b}
\end{array}$$

$$\begin{array}{c|c}
CH_2 \\
C \\
CH_2
\end{array}$$

$$\begin{array}{c|c}
CH_2 \\
C \\
CH_2
\end{array}$$

$$\begin{array}{c|c}
CH_2 \\
CH_2
\end{array}$$

(In the formula, R_{4h} , R_{4i} , R_{4j} , R_{4k} , R_{4l} , R_{4m} , and R_{4n} each independently represent SO₂R₄₀ (R₄₀ represents OH, a halogen atom, ONa, OK, or OR401. (R401 represents a linear or branched alkyl group having 1 to 8 carbon 5 atoms, or a substituted or unsubstituted phenyl group.)), a hydrogen atom, a halogen atom, an alkyl group having 1 to 20 carbon atoms, an alkoxy group having 1 to 20 carbon atoms, an OH group, an NH2 group, an NO2 group, COOR4p (R4p represents an H atom, 10 an Na atom, or a K atom.), an acetamide group, an OPh group, an NHPh group, a CF3 group, a C2F5 group, or a C_3F_7 group (Ph represents a phenyl group.), and at least one of these groups represents SO_2R_{40} . In addition, m represents an integer selected from 0 to 15 8. Zia represents a linear alkylene chain having 1 to 4 carbon atoms. The linear alkylene chain has at least one linear or branched alkyl group, or at least one alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a 20

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cyclohexyl structure at a terminal thereof. Z_{1b} represents a hydrogen atom, or a linear or branched alkyl group, aryl group, or aralkyl group which may be substituted by an aryl group. When multiple units exist, R_{4h} , R_{4i} , R_{4j} , R_{4k} , R_{4l} , R_{4m} , R_{4n} , R_{4o} , R_{4o1} , R_{4p} , Z_{1a} , Z_{1b} , and m each independently have the above meaning for each unit.)

Examples of the compound represented by the chemical formula (49A) or (49B) include: sulfonic 10 acids such as 1-naphthylamine-5-sulfonic acid, 1-naphthylamine-4-sulfonic acid, 1-naphthylamine-8-sulfonic acid, 2-naphthylamine-5-sulfonic acid, 1-naphthylamine-6-sulfonic acid, 1-naphthylamine-7-sulfonic acid, 1-naphthylamine-2-ethoxy-6-sulfonic acid, 1-amino-2-naphthol-4-sulfonic acid, 6-amino-1-naphthol-3-sulfonic acid, sodium 1-amino-8-naphthol-2,4-sulfonate, sodium 1-amino-8-naphthol-3,6-sulfonate; and alkali metal salts and esterified products of the sulfonic acids.

When A₃ represents a substituted or unsubstituted heterocyclic structure containing one or more of N, S, and O, A₃ may represent any one of a pyridine ring, a piperazine ring, a furan ring, and a thiol ring as a heterocyclic ring. Examples of such a compound include: sulfonic acids such as 2-aminopyridine-6-sulfonic acid and 2-aminopiperazine-6-sulfonic acid; and alkali metal salts and

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esterified products of the sulfonic acids.

As described above, examples of a group forming an ester bond with a sulfonic acid in the case of a sulfonate include a group containing a substituted or unsubstituted aliphatic hydrocarbon structure, a substituted or unsubstituted aromatic ring structure, and a substituted or unsubstituted heterocyclic structure. Furthermore, a linear or branched alkyl group having 1 to 8 carbon atoms, a substituted or unsubstituted phenyl group, or the like is preferable. From the viewpoint of, for example, ease of esterification, one having a group such as OCH₃, OC₂H₅, OC₆H₅, OC₃H₇, OC₄H₉, OCH(CH₃)₂, OCH₂C (CH₃)₃, or OC(CH₃)₃ is more preferable.

15 (Method of producing polyhydroxyalkanoate having unit represented by chemical formula (1))

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A reaction between a polyhydroxyalkanoate containing a unit represented by the chemical formula (29) and an aminosulfonic acid compound represented by the chemical formula (30) in the present invention will be described in detail.

The amount of the compound represented by the chemical formula (30) to be used in the present invention is in the range of 0.1 to 50.0 times mole, or preferably 1.0 to 20.0 times mole with respect to the unit represented by the chemical formula (29) to be used as a starting material. An example of a

method of producing an amide bond from a carboxylic acid and an amine in the present invention includes a condensation reaction by virtue of heat dehydration. In particular, from the viewpoint of achieving a mild reaction condition under which an ester bond of a 5 polymer main chain is not cleaved, a method is effective, which involves: activating a carboxylic acid portion with an activator to produce an active acyl intermediate; and allowing the intermediate to react with an amine. Examples of the active acyl 10 intermediate include an acid halide, an acid anhydride, and an active ester. In particular, a method of forming an amide bond in an identical reaction field by using a condensing agent is preferable from the viewpoint of simplifying a 15 production process.

If required, the active acyl intermediate may be isolated as an acid halide before being subjected to a condensation reaction with an amine.

A phosphoric acid-based condensing agent used for polycondensation of an aromatic polyamide, a carbodiimide-based condensing agent used for synthesizing a peptide, an acid chloride-based condensing agent, or the like can be appropriately selected as a condensing agent to be used depending 25 on the combination of the chemical formulae (30) and (29).

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Examples of the phosphoric acid-based condensing agent include a phosphite-based condensing agent, a phosphorus chloride-based condensing agent, a phosphoric anhydride-based condensing agent, a phosphate-based condensing agent, and a phosphoric amide-based condensing agent.

A phosphite-based condensing agent or the like can be used in the reaction of the present invention. Examples of a phosphite used at this time include triphenyl phosphite, diphenyl phosphite, tri-o-tolyl 10 phosphite, di-o-tolyl phosphite, tri-m-tolyl phosphite, di-m-tolyl phosphite, tri-p-tolyl phosphite, di-p-tolyl phosphite, di-o-chlorophenyl phosphite, tri-p-chlorophenyl phosphite, di-pchlorophenyl phosphite, trimethyl phosphite, and 15 triethyl phosphite. Of those, triphenyl phosphite is preferably used. A metal salt such as lithium chloride or calcium chloride may be added for improving the solubility, reactivity, and the like of 20 a polymer.

Examples of the carbodiimide-based condensing agent include dicyclohexyl carbodiimide (DCC), N-ethyl-N'-3-dimethylaminopropyl carbodiimide (EDC=WSCI), and diisopropyl carbodiimide (DIPC). DCC or WSCI may be used in combination with N-hydroxysuccinimide (HONSu), 1-hydroxybenzotriazole (HOBt), 3-hydroxy-4-oxo-3,4-dihydro-1,2,3-

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benzotriazine (HOObt), or the like.

The amount of the condensing agent to be used is in the range of 0.1 to 50 times mole, or preferably 1 to 20 times mole with respect to the compound represented by the chemical formula (29).

A solvent may be used as required in the reaction of the present invention. Examples of an available solvent include: hydrocarbons such as hexane, cyclohexane, and heptane; ketones such as acetone and methyl ethyl ketone; ethers such as dimethyl ether, diethyl ether, and tetrahydrofuran; halogenated hydrocarbons such as dichloromethane, chloroform, carbon tetrachloride, dichloroethane, and trichloroethane; aromatic hydrocarbons such as benzene and toluene; aprotic polar solvents such as N, N-dimethylformamide, dimethyl sulfoxide, dimethyl acetamide, and hexamethylphosphoramide; pyridine derivatives such as pyridine and picoline; and Nmethylpyrrolidone. Pyridine, N-methylpyrrolidone, or the like is particularly preferably used. The amount of the solvent to be used can be appropriately determined in accordance with kinds of a starting material and a base, a reaction condition, and the like.

25 A reaction temperature is not particularly limited in the method of the present invention, but is generally in the range of -20°C to the boiling

point of a solvent. However, it is preferable to perform the reaction at an optimum temperature suited for a condensing agent to be used.

In the method of the present invention, a reaction time is generally in the range of 1 to 48 hours. The reaction time is particularly preferably in the range of 1 to 10 hours.

A thus produced reaction solution containing a

polyhydroxyalkanoate having a unit represented by the chemical formula (1) in the present invention can be 10 collected and purified by, for example, distillation as an ordinary method. Alternatively, the reaction solution can be collected by mixing a solvent (for example, water, an alcohol such as methanol or ethanol, or an ether such as dimethyl ether, diethyl 15 ether, or tetrahydrofuran) evenly with the reaction solution; and reprecipitating a target polyhydroxyalkanoate having a unit represented by the chemical formula (1). The resultant polyhydroxyalkanoate having a unit represented by the 20 chemical formula (1) can be subjected to isolation purification as required. A method for the isolation purification is-not-particularly limited, and a method involving reprecipitation using a solvent insoluble in the polyhydroxyalkanoate represented by 25 the chemical formula (1), a method according to column chromatography, dialysis, or the like can be

used.

When an R portion in the chemical formula (1) is $-A_1-SO_3H$, a method can be adopted as another production method of the present invention, which involves methyl esterifying the R portion in the 5 chemical formula (1) into -A₁-SO₃CH₃- using a methylesterifying agent after a condensation reaction with an amine. Examples of an available methylesterifying agent include those used for methyl esterification of an aliphatic acid in gas 10 chromatography. Examples of a methyl esterification method include: acid catalyst methods such as a hydrochloric acid-methanol method, a boron trifluoride-methanol method, and a sulfuric acidmethanol method; and base catalyst methods such as a 15 sodium methoxide method, a tetramethylguanidine method, and a trimethylsilyldiazomethane method. those, a trimethylsilyldiazomethane method is preferable because methylation can be performed under a moderate condition. 20

Examples of a solvent to be used in the reaction of the present invention include: hydrocarbons such as hexane, cyclohexane, and heptane; alcohols such as methanol and ethanol; halogenated hydrocarbons such as dichloromethane, chloroform, carbon tetrachloride, dichloroethane, and trichloroethane; and aromatic hydrocarbons such as

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benzene and toluene. Halogenated hydrocarbons and the like are particularly preferably used. The amount of the solvent to be used can be appropriately determined in accordance with a starting material, a reaction condition, and the like. A reaction temperature is not particularly limited in the method of the present invention, but is generally in the range of -20°C to 30°C. However, it is preferable to perform the reaction at an optimum temperature suited for a condensing agent and a reagent to be used.

In addition, in the present invention, a polyhydroxyalkanoate containing a unit represented by the chemical formula (38) can be produced through the steps of: allowing a polyhydroxyalkanoate having a unit represented by the chemical formula (36) to react with a base; and allowing the compound obtained in the foregoing step to react with a compound represented by the chemical formula (37).

$$\begin{array}{c|c}
 & H \\
\hline
 & Z_{36a} - O \\
\hline
 & Z_{36b}
\end{array}$$
(36)

20 (In the formula, Z_{36a} represents a linear alkylene chain having 1 to 4 carbon atoms. The linear alkylene chain has at least one linear or branched alkyl group, or at least one alkyl group containing a residue having any one of a phenyl structure, a

25 thienyl structure, and a cyclohexyl structure at a

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terminal thereof. Z_{36b} represents a hydrogen atom, or a linear or branched alkyl group, aryl group, or aralkyl group which may be substituted by an aryl group. When multiple units exist, Z_{36a} and Z_{36b} each independently have the above meaning for each unit.)

(In the formula, R_{37} represents $-A_{37}-SO_2R_{37a}$. R_{37a} represents OH, a halogen atom, ONa, OK, or OR_{37b} . R_{37b} and A_{37} are each independently selected from groups each having a substituted or unsubstituted aliphatic hydrocarbon structure, a substituted or unsubstituted aromatic ring structure, or a substituted or unsubstituted or unsubstituted heterocyclic structure. When multiple units exist, R_{37} , R_{37a} , R_{37b} , and A_{37} each independently have the above meaning for each unit.)

(In the formula, R_{38} represents $-A_{38}-SO_2R_{38a}$. R_{38a} represents OH, a halogen atom, ONa, OK, or OR_{38b} . R_{38b} and A_{38} each independently represent a group having a substituted or unsubstituted aliphatic hydrocarbon structure, a substituted or unsubstituted aromatic

ring structure, or a substituted or unsubstituted heterocyclic structure. Z_{38a} represents a linear alkylene chain having 1 to 4 carbon atoms. The linear alkylene chain has at least one linear or branched alkyl group, or at least one alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof. Z_{38b} represents a hydrogen atom, or a linear or branched alkyl group, aryl group, or aralkyl group which may be substituted by an aryl group. When multiple units exist, R₃₈, R_{38a}, R_{38b}, A₃₈, Z_{38a}, and Z_{38b} each independently have the above meaning for each unit.)

More specifically, in the compound represented

by the chemical formulae (36) and (38) to be used in
the present invention, Z_{36a} and Z_{38a} each represent a
linear alkylene chain having 1 to 4 carbon atoms.

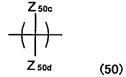
The linear alkylene chain is arbitrarily substituted
by at least one linear or branched alkyl group, or at
least one alkyl group containing a residue having any
one of a phenyl structure, a thienyl structure, and a
cyclohexyl structure at a terminal thereof. The
linear alkylene chain structure represented by each
of Z_{36a} and Z_{38a} is preferably selected from the
following (A) to (D).

(A) When the linear alkylene chain has 1 carbon atom, in the linear alkylene chain structure

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represented by the chemical formula (50), one of Z_{50c} and Z_{50d} represents a linear or branched alkyl group, or an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof.



(B) When the linear alkylene chain has 2 carbon atoms, in the linear alkylene chain structure represented by the chemical formula (51), one of Z_{51c} , Z_{51d} , Z_{51e} , and Z_{51f} represents a linear or branched alkyl group, or an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof.



(C) When the linear alkylene chain has 3 carbon atoms, in the linear alkylene chain structure represented by the chemical formula (52), one of Z_{52c} , Z_{52d} , Z_{52e} , Z_{52f} , Z_{52g} , and Z_{52h} represents a linear or branched alkyl group, or an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof.

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(D) When the linear alkylene chain has 4 carbon atoms, in the linear alkylene chain structure represented by the chemical formula (53), one of Z_{53c} , Z_{53d} , Z_{53e} , Z_{53f} , Z_{53g} , Z_{53h} , Z_{53i} , and Z_{53j} represents a linear or branched alkyl group, or an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof.

When a substituent selected from Z_{50c} , Z_{50d} , Z_{51c} , Z_{51d} , Z_{51e} , Z_{51e} , Z_{52c} , Z_{52d} , Z_{52e} , Z_{52f} , Z_{52g} , Z_{52h} , Z_{53c} , Z_{53d} , Z_{53e} , Z_{53f} , Z_{53g} , Z_{53h} , Z_{53i} , and Z_{53j} described in the chemical formulae (50), (51), (52), and (53)

represents an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof, the substituent is more preferably selected from substituents represented by the chemical formulae (14), (15), (16), and (17).

$$--(CH_2)k_{14} - CH_3$$
 (14)

(In the formula, k_{14} represents an integer selected from 0 to 8. When multiple units exist, k_{14} 's each

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independently have the above meaning for each unit.)

$$--(CH2)k15---(CH3) CH3 (15)$$

(In the formula, k_{15} represents an integer selected from 0 to 7. When multiple units exist, k_{15} 's each independently have the above meaning for each unit.) $-(CH_2)k_{\overline{16}}R_{16}$ (16)

(In the formula, k_{16} represents an integer selected from 1 to 8. R_{16} represents a substituent containing a residue having any one of a phenyl structure and a thienyl structure. When multiple units exist, k_{16} and R_{16} each independently have the above meaning for each unit.)

$$-(CH_2)k_{17}$$

(In the formula, R_{17} represents a substituent to a cyclohexyl group selected from an H atom, a CN group, an NO_2 group, a halogen atom, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a CF_3 group, a C_2F_5 group, and a C_3F_7 group. k_{17} represents an integer selected from 0 to 8. When multiple units exist, k_{17} and R_{17} each independently have the above meaning for each unit.)

In addition, R_{16} in the chemical formula (16), that is, a residue having any one of a phenyl structure and a thienyl structure is selected from the group of residues represented by the chemical formulae (18), (19), (20), (21), (22), (23), (24),

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(25), (26), (27), and (28).

Here, the chemical formula (18) represents a group of unsubstituted or substituted phenyl groups.

(In the formula, R₁₈ represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO₂ group, a CH₃ group, a C₂H₅ group, a C₃H₇ group, a CH=CH₂ group, COOR_{18a} (R_{18a} represents an H atom, an Na atom, or a K atom.), a CF₃ group, a C₂F₅ group, and a C₃F₇ group. When multiple units exist, R₁₈'s may be different for each unit.)

The chemical formula (19) represents a group of unsubstituted or substituted phenoxy groups.

(In the formula, R₁₉ represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO₂ group, a CH₃ group, a C₂H₅ group, a C₃H₇ group, a SCH₃ group, a CF₃ group, a C₂F₅ group, and a C₃F₇ group. When multiple units exist, R₁₉'s may be different for each unit.)

The chemical formula (20) represents a group of unsubstituted or substituted benzoyl groups.

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(In the formula, R_{20} represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a CF_3 group, a C_2F_5 group, and a C_3F_7 group. When multiple units exist, R_{20} 's may be different for each unit.)

The chemical formula (21) represents a group of unsubstituted or substituted phenylsulfanyl groups.

(In the formula, R_{21} represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO_2 group, $COOR_{21a}$, SO_2R_{21b} (R_{21a} represents H, Na, K, CH_3 , or C_2H_5 , and R_{21b} represents OH, ONa, OK, a halogen atom, OCH₃, or OC_2H_5 .), a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(CH_3)_2$ -CH group, and a $(CH_3)_3$ -C group. When multiple units exist, R_{21} 's may be different for each unit.)

The chemical formula (22) represents a group of unsubstituted or substituted (phenylmethyl)sulfanyl groups.

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$$R_{22}$$
 CH_2 CS (22)

(In the formula, R_{22} represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO_2 group, $COOR_{22a}$, SO_2R_{22b} (R_{22a} represents H, Na, K, CH_3 , or C_2H_5 , and R_{22b} represents OH, ONa, OK, a halogen atom, OCH₃, or OC_2H_5 .), a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(CH_3)_2$ -CH group, and a $(CH_3)_3$ -C group. When multiple units exist, R_{22} 's may be different for each unit.)

The chemical formula (23) represents a 2-thienyl group.

The chemical formula (24) represents a 2-thienylsulfanyl group.

The chemical formula (25) represents a 2-thienylcarbonyl group.

The chemical formula (26) represents a group of unsubstituted or substituted phenylsulfinyl groups.

(In the formula, R26 represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO₂ group, COOR_{26a}, SO₂R_{26b} (R_{26a} 5 represents H, Na, K, CH₃, or C₂H₅, and R_{26b} represents OH, ONa, OK, a halogen atom, OCH₃, or OC₂H₅.), a CH₃ group, a C₂H₅ group, a C₃H₇ group, a (CH₃)₂-CH group, and a $(CH_3)_3$ -C group. When multiple units exist, R_{26} 's may be different for each unit.)

10 The chemical formula (27) represents a group of unsubstituted or substituted phenylsulfonyl groups.

(27)

(In the formula, R_{27} represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO₂ group, COOR_{27a}, SO₂R_{27b} (R_{27a} 15 represents H, Na, K, CH_3 , or C_2H_5 , and R_{27b} represents OH, ONa, OK, a halogen atom, OCH₃, or OC₂H₅.), a CH₃ group, a C₂H₅ group, a C₃H₇ group, a (CH₃)₂-CH group, and a (CH₃)₃-C group. When multiple units exist, R₂₇'s may be different for each unit.) 20

The chemical formula (28) represents a (phenylmethyl) oxy group.

On the other hand, examples of the compound represented by the chemical formula (37) to be used in the present invention include 2-acrylamide-2-methylpropanesulfonic acid, and alkali metal salts and esterified products thereof.

(Method of producing polyhydroxyalkanoate represented by chemical formula (38))

A reaction between the polyhydroxyalkanoate

10 containing a unit represented by the chemical formula

(36) and the compound represented by the chemical

formula (37) in the present invention will be

described in detail.

The present invention can be achieved by

15 subjecting a α-methylene or a α-methine adjacent to
a carbonyl group in a polymer main chain to a Michael
addition reaction with the compound represented by
the chemical formula (37). To be specific, the
present invention can be achieved by allowing the

20 polyhydroxyalkanoate containing a unit represented by
the chemical formula (36) to react with a base
capable of forming a α-methylene or a α-methine,
which is adjacent to a carbonyl group in the polymer
main chain of the polyhydroxyalkanoate containing a

25 unit represented by the chemical formula (36), into

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an anion under a Michael addition reaction condition; and allowing the resultant to react with the compound represented by the chemical formula (37). In the present invention, the amount of the compound represented by the chemical formula (37) to be used is 0.001 to 100 times mole, or preferably 0.01 to 10 times mole with respect to the unit represented by the chemical formula (36).

A solvent to be used in the reaction of the

present invention is not particularly limited as long
as it is inactive to the reaction and dissolves the
staring material to some extent. Examples of such a
solvent include: aliphatic hydrocarbons such as
hexane, cyclohexane, heptane, ligroin, and petroleum

ther; aromatic hydrocarbons such as benzene, toluene,
and xylene; ethers such as diethyl ether, diisopropyl
ether, tetrahydrofuran, dioxane, dimethoxyethane, and
diethyleneglycoldimethylether; and amides such as
formamide, N,N-dimethylformamide, N,N
dimethylacetamide, N-methyl-2-pyrrolidone, N-

The reaction is performed in the presence of a base. Examples of a base to be used include: lithium alkyls such as methyl lithium and butyl lithium; alkali metal disilazides such as lithium hexamethyl disilazide, sodium hexamethyl disilazide, and

methylpyrrolidinone, and hexamethylphosphortriamide.

Of those, tetrahydrofuran is preferable.

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potassium hexamethyl disilazide; and lithium amides such as lithium diisopropylamide and lithium dicyclohexylamide. Of those, lithium diisopropylamide is preferable. In addition, the amount of the base to be used is 0.001 to 100 times mole, or preferably 0.01 to 10 times mole with respect to the unit represented by the chemical formula (36).

In the method of the present invention, a

reaction temperature is generally in the range of

-78°C to 40°C, or preferably in the range of -78°C to

30°C.

In the method of the present invention, a reaction time is generally in the range of 10 minutes to 24 hours. The reaction time is particularly preferably in the range of 10 minutes to 4 hours.

In addition, in the polyhydroxyalkanoate having a unit represented by the chemical formula (5) of the present invention, the polyhydroxyalkanoate having a unit represented by the chemical formula (32) can be produced by hydrolyzing a side chain ester portion of a polyhydroxyalkanoate having a unit represented by the chemical formula (31) as a starting material in the presence of an acid or an alkali, or by subjecting the polyhydroxyalkanoate to hydrogenolysis including catalytic reduction.

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(In the formula, R₃₂ represents hydrogen or a group for forming a salt. Z_{32a} represents a linear alkylene chain having 1 to 4 carbon atoms. The linear alkylene chain has at least one linear or branched alkyl group, or at least one alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof. Z_{32b} represents a hydrogen atom, or a linear or branched alkyl group, aryl group, or aralkyl group which may be substituted by an aryl group. m represents an integer selected from 0 to 8. When multiple units exist, R₃₂, Z_{32a}, Z_{32b}, and m each independently have the above meaning for each unit.)

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(In the formula, R_{31} represents a linear or branched alkyl or aralkyl group having 1 to 12 carbon atoms. Z_{31a} represents a linear alkylene chain having 1 to 4 carbon atoms. The linear alkylene chain has at least one linear or branched alkyl group, or at least one alkyl group containing a residue having any one of a

phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof. Z_{31b} represents a hydrogen atom, or a linear or branched alkyl group, aryl group, or aralkyl group which may be substituted by an aryl group. m represents an integer selected from 0 to 8. When multiple units exist, R_{31} , Z_{31a} , Z_{31b} , and m each independently have the above meaning for each unit.)

More specifically, in each of the compounds

represented by the chemical formulae (31) and (32) to
be used in the present invention, Z_{31a} and Z_{32a} each
represent a linear alkylene chain having 1 to 4
carbon atoms. The linear alkylene chain is
arbitrarily substituted by at least one linear or

branched alkyl group, or at least one alkyl group
containing a residue having any one of a phenyl
structure, a thienyl structure, and a cyclohexyl
structure at a terminal thereof. The linear alkylene
chain structure represented by each of Z_{31a} and Z_{32a} is

preferably selected from the following (A) to (D).

(A) When the linear alkylene chain has 1 carbon atom, in the linear alkylene chain structure represented by the chemical formula (54), one of Z_{54c} and Z_{54d} represents a linear or branched alkyl group, or an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof.

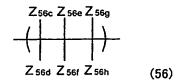
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(B) When the linear alkylene chain has 2 carbon atoms, in the linear alkylene chain structure represented by the chemical formula (55), one of Z_{55c} , Z_{55d} , Z_{55e} , and Z_{55f} represents a linear or branched alkyl group, or an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof.

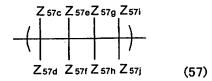


(C) When the linear alkylene chain has 3 carbon atoms, in the linear alkylene chain structure represented by the chemical formula (56), one of Z_{56c}, Z_{56d}, Z_{56e}, Z_{56f}, Z_{56g}, and Z_{56h} represents a linear or branched alkyl group, or an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof.



20 (D) When the linear alkylene chain has 4 carbon atoms, in the linear alkylene chain structure

represented by the chemical formula (57), one of Z_{57c} , Z_{57d} , Z_{57e} , Z_{57f} , Z_{57g} , Z_{57h} , Z_{57i} , and Z_{57j} represents a linear or branched alkyl group, or an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof.



When a substituent selected from Z_{54c}, Z_{54d}, Z_{55c}, Z_{55d} , Z_{55e} , Z_{55f} , Z_{56c} , Z_{56d} , Z_{56e} , Z_{56f} , Z_{56g} , Z_{56h} , Z_{57c} , Z_{57d} , Z_{57e} , Z_{57f} , Z_{57g} , Z_{57h} , Z_{57i} , and Z_{57j} described in the 10 chemical formulae (54), (55), (56), and (57) represents a linear or branched alkyl group, or an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof, the 15 substituent is more preferably selected from substituents represented by the chemical formulae (14), (15), (16), and (17). ---(CH₂)k₁₄--CH₃

$$--(CH_2)k_{14} - CH_3$$
 (14)

(In the formula, k_{14} represents an integer selected 20 from 0 to 8. When_multiple_units exist, k14's each independently have the above meaning for each unit.)

$$--(CH_2)n_{15}$$
 CH_3 (15)

(In the formula, k_{15} represents an integer selected

from 0 to 7. When multiple units exist, k_{15} 's each independently have the above meaning for each unit.) $-(CH_2)k_{16}-R_{16}$ (16)

(In the formula, k_{16} represents an integer selected from 1 to 8. R_{16} represents a substituent containing a residue having any one of a phenyl structure and a thienyl structure. When multiple units exist, k_{16} and R_{16} each independently have the above meaning for each unit.)

$$-(CH_2)k_{17}$$

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(In the formula, R_{17} represents a substituent to a cyclohexyl group selected from an H atom, a CN group, an NO_2 group, a halogen atom, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a CF_3 group, a C_2F_5 group, and a C_3F_7 group. k_{17} represents an integer selected from 0 to 8. When multiple units exist, k_{17} and R_{17} each independently have the above meaning for each unit.)

In addition, R_{16} in the chemical formula (16), that is, a residue having any one of a phenyl structure and a thienyl structure is selected from the group of residues represented by the chemical formulae (18), (19), (20), (21), (22), (23), (24), (25), (26), (27), and (28).

Here, the chemical formula (18) represents a group of unsubstituted or substituted phenyl groups.

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(In the formula, R_{18} represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $CH=CH_2$ group, $COOR_{18a}$ (R_{18a} represents an H atom, an Na atom, or a K atom.), a CF_3 group, a C_2F_5 group, and a C_3F_7 group. When multiple units exist, R_{18} 's may be different for each unit.)

The chemical formula (19) represents a group of unsubstituted or substituted phenoxy groups.

(In the formula, R_{19} represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a SCH_3 group, a CF_3 group, a C_2F_5 group, and a C_3F_7 group. When multiple units exist, R_{19} 's may be different for each unit.)

The chemical formula (20) represents a group of unsubstituted or substituted benzoyl groups.

(In the formula, R20 represents a substituent to an

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aromatic ring selected from an H atom, a halogen atom, a CN group, an NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a CF_3 group, a C_2F_5 group, and a C_3F_7 group. When multiple units exist, R_{20} 's may be different for each unit.)

The chemical formula (21) represents a group of unsubstituted or substituted phenylsulfanyl groups.

(In the formula, R₂₁ represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO₂ group, COOR_{21a}, SO₂R_{21b} (R_{21a} represents H, Na, K, CH₃, or C₂H₅, and R_{21b} represents OH, ONa, OK, a halogen atom, OCH₃, or OC₂H₅.), a CH₃ group, a C₂H₅ group, a C₃H₇ group, a (CH₃)₂-CH group, and a (CH₃)₃-C group. When multiple units exist, R₂₁'s may be different for each unit.)

The chemical formula (22) represents a group of unsubstituted or substituted (phenylmethyl)sulfanyl groups.

$$R_{22}$$
 CH_2 CS

(In the formula, R_{22} represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO_2 group, $COOR_{22a}$, SO_2R_{22b} (R_{22a}

represents H, Na, K, CH_3 , or C_2H_5 , and R_{22b} represents OH, ONa, OK, a halogen atom, OCH₃, or OC₂H₅.), a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(CH_3)_2$ -CH group, and a $(CH_3)_3$ -C group. When multiple units exist, R_{22} 's may be different for each unit.)

The chemical formula (23) represents a 2-thienyl group.

The chemical formula (24) represents a 2-10 thienylsulfanyl group.

The chemical formula (25) represents a 2-thienylcarbonyl group.

The chemical formula (26) represents a group of unsubstituted or substituted phenylsulfinyl groups.

(In the formula, R₂₆ represents a substituent to an aromatic ring selected from an H atom, a halogen atom, 20 a CN group, an NO₂ group, COOR_{26a}, SO₂R_{26b} (R_{26a} represents H, Na, K, CH₃, or C₂H₅, and R_{26b} represents OH, ONa, OK, a halogen atom, OCH₃, or OC₂H₅.), a CH₃

group, a C_2H_5 group, a C_3H_7 group, a $(CH_3)_2$ -CH group, and a $(CH_3)_3$ -C group. When multiple units exist, R_{26} 's may be different for each unit.)

The chemical formula (27) represents a group of unsubstituted or substituted phenylsulfonyl groups.

(In the formula, R₂₇ represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO₂ group, COOR_{27a}, SO₂R_{27b} (R_{27a}

10 represents H, Na, K, CH₃, or C₂H₅, and R_{27b} represents OH, ONa, OK, a halogen atom, OCH₃, or OC₂H₅.), a CH₃ group, a C₂H₅ group, a C₃H₇ group, a (CH₃)₂-CH group, and a (CH₃)₃-C group. When multiple units exist, R₂₇'s may be different for each unit.)

The chemical formula (28) represents a (phenylmethyl)oxy group.

(Method of producing polyhydroxyalkanoate represented by chemical formula (32))

Detailed description will be given of a method of producing the polyhydroxyalkanoate having a unit represented by the chemical formula (32) by hydrolyzing a side chain ester portion of a

polyhydroxyalkanoate having a unit represented by the chemical formula (31) in the presence of an acid or an alkali, or by subjecting the polyhydroxyalkanoate to hydrogenolysis including catalytic reduction in the present invention.

In the case where hydrolysis in the presence of an acid or an alkali is employed, the hydrolysis can be performed by using, in an aqueous solution or a hydrophilic organic solvent such as methanol, ethanol, 10 tetrahydrofuran, dioxane, dimethylformamide, or dimethyl sulfoxide as a solvent, an aqueous solution of an inorganic acid such as hydrochloric acid, sulfuric acid, nitric acid, or phosphoric acid, an organic acid such as trifluoroacetic acid, trichloroacetic acid, p-toluenesulfonic acid, or 15 methanesulfonic acid, an aqueous caustic alkali such as sodium hydroxide or potassium hydroxide, an aqueous solution of an alkali carbonate such as sodium carbonate or potassium carbonate, or an alcohol solution of a metal alkoxide such as sodium 20 methoxide or sodium ethoxide. A reaction temperature is generally in the range of 0°C to 40°C, or preferably in the range of 0°C to 30°C. A reaction time is generally in the range of 0.5 to 48 hours. When hydrolysis is performed in the presence of an 25 acid or an alkali, in each case, an ester bond of a main chain is also cleaved, and a reduction in

molecular weight is observed in some cases.

A method of obtaining a carboxylic acid by way of hydrogenolysis including catalytic reduction is performed as follows. That is, in an appropriate solvent, in the temperature range of -20°C to the 5 boiling point of the solvent used, or preferably 0 to 50°C, in the presence of a reduction catalyst, hydrogen is allowed to act under normal or increased pressure to perform catalytic reduction. Examples of the solvent used include water, methanol, ethanol, 10 propanol, hexafluoroisopropanol, ethyl acetate, diethyl ether, tetrahydrofuran, dioxane, benzene, toluene, dimethylformamide, pyridine, and Nmethylpyrrolidone. A mixed solvent of the above 15 solvents may also be used. A catalyst such as palladium, platinum, or rhodium which is used singly or used while being carried by a carrier, Raney nickel, or the like is used as the reduction catalyst. A reaction time is generally in the range of 0.5 to 72 hours. A thus produced reaction solution 20 containing a polyhydroxyalkanoate having a unit represented by the chemical formula (32) is collected as a crude polymer by: removing the catalyst through filtration; and removing the solvent through distillation or the like. The resultant 25 polyhydroxyalkanoate having a unit represented by the

chemical formula (32) can be subjected to isolation

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purification as required. A method for the isolation purification is not particularly limited, and a method involving reprecipitation using a solvent insoluble in the polyhydroxyalkanoate having a unit represented by the chemical formula (32), a method according to column chromatography, dialysis, or the like can be used. Provided, however, that even in the case where catalytic reduction is employed, an ester bond of a main chain is cleaved, and a reduction in molecular weight is observed in some cases.

In addition, in the polyhydroxyalkanoate having a unit represented by the chemical formula (5) of the present invention, a polyhydroxyalkanoate having a unit represented by the chemical formula (58) can be produced by esterifying a polyhydroxyalkanoate represented by the chemical formula (59) as a staring material by means of an esterifying agent.

$$COOR_{58}$$
 $(CH_2)m$
 $Z_{58a}-O$
 Z_{58b}
 (58)

20 (In the formula, R_{58} represents a linear or branched alkyl or aralkyl group having 1 to 12 carbon atoms. Z_{58a} represents a linear alkylene chain having 1 to 4 carbon atoms. The linear alkylene chain has at least one linear or branched alkyl group, or at least one

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alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof. Z_{58b} represents a hydrogen atom, or a linear or branched alkyl group, aryl group, or aralkyl group which may be substituted by an aryl group. m represents an integer selected from 0 to 8. When multiple units exist, R_{58} , Z_{58a} , Z_{58b} , and m each independently have the above meaning for each unit.)

(In the formula, R₅₉ represents hydrogen or a group for forming a salt. Z_{59a} represents a linear alkylene chain having 1 to 4 carbon atoms. The linear alkylene chain has at least one linear or branched alkyl group, or at least one alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof. Z_{59b} represents a hydrogen atom, or a linear or branched alkyl group, aryl group, or 20 aralkyl group which may be substituted by an aryl group. m represents an integer selected from 0 to 8. When multiple units exist, R₅₉, Z_{59a}, Z_{59b}, and m each

More specifically, in each of the compounds

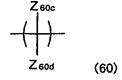
independently have the above meaning for each unit.)

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represented by the chemical formulae (58) and (59) to be used in the present invention, Z_{58a} and Z_{59a} each represent a linear alkylene chain having 1 to 4 carbon atoms. The linear alkylene chain is

5 arbitrarily substituted by at least one linear or branched alkyl group, or at least one alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof. The linear alkylene chain structure represented by each of Z_{58a} and Z_{59a} is preferably selected from the following (A) to (D).

(A) When the linear alkylene chain has 1 carbon atom, in the linear alkylene chain structure represented by the chemical formula (60), one of Z_{60c} and Z_{60d} represents a linear or branched alkyl group, or an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof.



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(B) When the linear alkylene chain has 2 carbon atoms, in the linear alkylene chain structure represented by the chemical formula (61), one of Z_{61c} , Z_{61d} , Z_{61e} , and Z_{61f} represents a linear or branched alkyl group, or an alkyl group containing a residue having any one of

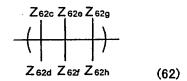
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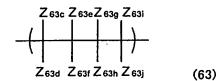
a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof.



(C) When the linear alkylene chain has 3 carbon atoms, in the linear alkylene chain structure represented by the chemical formula (62), one of Z_{62c} , Z_{62d} , Z_{62e} , Z_{62f} , Z_{62g} , and Z_{62h} represents a linear or branched alkyl group, or an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof.



(D) When the linear alkylene chain has 4 carbon atoms, in the linear alkylene chain structure represented by the chemical formula (63), one of Z_{63c}, Z_{63d}, Z_{63e}, Z_{63f},
15 Z_{63g}, Z_{63h}, Z_{63i}, and Z_{63j} represents a linear or branched alkyl group, or an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof.



When a substituent selected from Z_{60c}, Z_{60d}, Z_{61c},

Z_{61d}, Z_{61e}, Z_{61f}, Z_{62c}, Z_{62d}, Z_{62e}, Z_{62f}, Z_{62g}, Z_{62h}, Z_{63c}, Z_{63d}, Z_{63e}, Z_{63f}, Z_{63g}, Z_{63h}, Z_{63i}, and Z_{63j} described in the chemical formulae (60), (61), (62), and (63) represents a linear or branched alkyl group, or an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof, the substituent is more preferably selected from substituents represented by the chemical formulae (14), (15), (16), and (17).

10 (14), (15), (16), and (17) $--(CH_2)k_{14}-CH_3$ (14)

(In the formula, k_{14} represents an integer selected from 0 to 8. When multiple units exist, k_{14} 's each independently have the above meaning for each unit.)

$$--(CH2)n15---CH3$$

$$CH3$$

$$(15)$$

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(In the formula, k_{15} represents an integer selected from 0 to 7. When multiple units exist, k_{15} 's each independently have the above meaning for each unit.) $-(CH_2)n_{16}-R_{16}$ (16)

20 (In the formula, k_{16} represents an integer selected from 1 to 8. R_{16} represents a substituent containing a residue having—any one—of a phenyl structure and a thienyl structure. When multiple units exist, k_{16} and R_{16} each independently have the above meaning for each unit.)

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$$-(CH_2)k_{17}$$

(In the formula, R_{17} represents a substituent to a cyclohexyl group selected from an H atom, a CN group, an NO_2 group, a halogen atom, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a CF_3 group, a C_2F_5 group, and a C_3F_7 group. k_{17} represents an integer selected from 0 to 8. When multiple units exist, k_{17} and R_{17} each independently have the above meaning for each unit.)

In addition, R₁₆ in the chemical formula (16), that is, a residue having any one of a phenyl structure and a thienyl structure is selected from the group of residues represented by the chemical formulae (18), (19), (20), (21), (22), (23), (24), (25), (26), (27), and (28).

Here, the chemical formula (18) represents a group of unsubstituted or substituted phenyl groups.

(In the formula, R₁₈ represents a substituent to an aromatic ring selected from an H atom, a halogen atom, 20 a CN group, an NO₂ group, a CH₃ group, a C₂H₅ group, a C₃H₇ group, a CH=CH₂ group, COOR_{18a} (R_{18a} represents an H atom, an Na atom, or a K atom.), a CF₃ group, a C₂F₅ group, and a C₃F₇ group. When multiple units exist, R₁₈'s may be different for each unit.)

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The chemical formula (19) represents a group of unsubstituted or substituted phenoxy groups.

(In the formula, R_{19} represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a SCH_3 group, a CF_3 group, a C_2F_5 group, and a C_3F_7 group. When multiple units exist, R_{19} 's may be different for each unit.)

The chemical formula (20) represents a group of unsubstituted or substituted benzoyl groups.

(In the formula, R_{20} represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a CF_3 group, a C_2F_5 group, and a C_3F_7 group. When multiple units exist, R_{20} 's may be different for each unit.)

The chemical formula (21) represents a group of unsubstituted or substituted phenylsulfanyl groups.

(In the formula, R_{21} represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO_2 group, $COOR_{21a}$, SO_2R_{21b} (R_{21a} represents H, Na, K, CH₃, or C_2H_5 , and R_{21b} represents OH, ONa, OK, a halogen atom, OCH₃, or OC_2H_5 .), a CH₃ group, a C_2H_5 group, a C_3H_7 group, a $(CH_3)_2$ -CH group, and a $(CH_3)_3$ -C group. When multiple units exist, R_{21} 's may be different for each unit.)

The chemical formula (22) represents a group of unsubstituted or substituted (phenylmethyl)sulfanyl groups.

(In the formula, R₂₂ represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO₂ group, COOR_{22a}, SO₂R_{22b} (R_{22a} represents H, Na, K, CH₃, or C₂H₅, and R_{22b} represents OH, ONa, OK, a halogen atom, OCH₃, or OC₂H₅.), a CH₃ group, a C₂H₅ group, a C₃H₇ group, a (CH₃)₂-CH group, and a (CH₃)₃-C group. When multiple units exist, R₂₂'s may be different for each unit.)

The chemical formula (23) represents a 2-thienyl group.

The chemical formula (24) represents a 2-

thienylsulfanyl group.

The chemical formula (25) represents a 2-thienylcarbonyl group.

The chemical formula (26) represents a group of unsubstituted or substituted phenylsulfinyl groups.

(In the formula, R₂₆ represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO₂ group, COOR_{26a}, SO₂R_{26b} (R_{26a} represents H, Na, K, CH₃, or C₂H₅, and R_{26b} represents OH, ONa, OK, a halogen atom, OCH₃, or OC₂H₅.), a CH₃ group, a C₂H₅ group, a C₃H₇ group, a (CH₃)₂-CH group, and a (CH₃)₃-C group. When multiple units exist, R₂₆'s may be different for each unit.)

The chemical formula (27) represents a group of unsubstituted or substituted phenylsulfonyl groups.

20 (In the formula, R_{27} represents a substituent to an aromatic ring selected from an H atom, a halogen atom,

a CN group, an NO_2 group, $COOR_{27a}$, SO_2R_{27b} (R_{27a} represents H, Na, K, CH_3 , or C_2H_5 , and R_{27b} represents OH, ONa, OK, a halogen atom, OCH₃, or OC_2H_5 .), a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(CH_3)_2$ -CH group, and a $(CH_3)_3$ -C group. When multiple units exist, R_{27} 's may be different for each unit.)

The chemical formula (28) represents a (phenylmethyl)oxy group.

Examples of the esterifying agent to be used 10 include diazomethane and DMF dimethylacetals. For example, the polyhydroxyalkanoate having a unit represented by the chemical formula (59) easily reacts with trimethylsilyldiazomethane, DMF 15 dimethylacetal, DMF diethylacetal, DMF dipropylacetal, DMF diisopropylacetal, DMF-n-butylacetal, DMF-tertbutylacetal, DMF dineopentylacetal, or the like to produce a corresponding ester. Furthermore, the polyhydroxyalkanoate is allowed to react with any one of alcohols such as methanol, ethanol, propanol, 20 isopropyl alcohol, butyl alcohol, sec-butyl alcohol, tert-butyl alcohol, pentyl alcohol, neopentyl alcohol, hexyl alcohol, heptyl alcohol, octyl alcohol, nonyl alcohol, decyl alcohol, and lauryl alcohol, or any one of saccharides for introducing a sugar structure 25

such as D-glucose, D-fructose, and otherwise by using an acid catalyst or a condensing agent such as DCC to produce an esterified polyhydroxyalkanoate.

In addition, in the present invention, a

5 polyhydroxyalkanoate containing a unit represented by
the chemical formula (35) can be produced through the
steps of: allowing a polyhydroxyalkanoate having a
unit represented by the chemical formula (33) to
react with a base; and allowing the compound obtained

10 in the foregoing step to react with a compound
represented by the chemical formula (34).

$$\begin{array}{c|c}
 & H \\
\hline
 & Z_{33a} - O \\
\hline
 & Z_{33b}
\end{array}$$
(33)

(In the formula, Z_{33a} represents a linear alkylene chain having 1 to 4 carbon atoms. The linear

15 alkylene chain has at least one linear or branched alkyl group, or at least one alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof. Z_{33b} represents a hydrogen atom, or 20 a linear or branched alkyl group, aryl group, or aralkyl group which—may-be-substituted by an aryl group. When multiple units exist, Z_{33a} and Z_{33b} each independently have the above meaning for each unit.) X(CH₂)mCOOR₃₄ (34)

25 (In the formula, m represents an integer selected

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from 0 to 8. X represents a halogen atom. R_{34} represents a linear or branched alkyl or aralkyl group having 1 to 12 carbon atoms.)

COOR₃₅

$$(CH2)m$$

$$- \left(\begin{array}{c|c} & & \\ & & \\ \hline & & \\$$

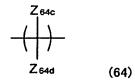
(In the formula, R₃₅ represents a linear or branched alkyl or aralkyl group having 1 to 12 carbon atoms. Z_{35a} represents a linear alkylene chain having 1 to 4 carbon atoms. The linear alkylene chain has at least one linear or branched alkyl group, or at least one alkyl group containing a residue having any one of a 10 phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof. Z35b represents a hydrogen atom, or a linear or branched alkyl group, aryl group, or aralkyl group which may be substituted by an aryl group. In addition, m 15 represents an integer selected from 0 to 8. When multiple units exist, R₃₅, Z_{35a}, Z_{35b}, and m each independently have the above meaning for each unit.)

Alternatively, the compound (33) can be produced via a ring-opening polymerizable cyclic compound.

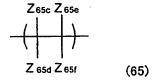
More specifically, in each of the compounds represented by the chemical formulae (33) and (35) in the present invention, Z_{33a} and Z_{35a} each represent a

linear alkylene chain having 1 to 4 carbon atoms. The linear alkylene chain is arbitrarily substituted by at least one linear or branched alkyl group, or at least one alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof. The linear alkylene chain structure represented by each of Z_{33a} and Z_{35a} is preferably selected from the following (A) to (D).

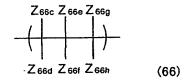
10 (A) When the linear alkylene chain has 1 carbon atom, in the linear alkylene chain structure represented by the chemical formula (33), one of Z_{64c} and Z_{64d} represents a linear or branched alkyl group, or an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof.



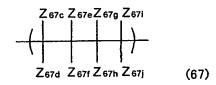
(B) When the linear alkylene chain has 2 carbon atoms, in the linear alkylene chain structure represented by the chemical formula (65), one of Z_{65c}, Z_{65d}, Z_{65e}, and Z_{65f} represents a linear or branched alkyl group, or an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof.



(C) When the linear alkylene chain has 3 carbon atoms, in the linear alkylene chain structure represented by the chemical formula (66), one of Z_{66c} , Z_{66d} , Z_{66e} , Z_{66e} , Z_{66e} , and Z_{66n} represents a linear or branched alkyl group, or an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof.



10 (D) When the linear alkylene chain has 4 carbon atoms, in the linear alkylene chain structure represented by the chemical formula (67), one of Z_{67c} , Z_{67d} , Z_{67e} , Z_{67f} , Z_{67g} , Z_{67h} , Z_{67i} , and Z_{67j} represents a linear or branched alkyl group, or an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof.



When a substituent selected from Z_{64c} , Z_{64d} , Z_{65c} , Z_{65d} , Z_{65e} , Z_{65e} , Z_{66c} , Z_{66d} , Z_{66e} , Z_{66f} , Z_{66g} , Z_{66h} , Z_{67c} , Z_{67d} , Z_{67e} , Z_{67f} , Z_{67g} , Z_{67h} , Z_{67i} , and Z_{67j} described in the

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chemical formulae (64), (65), (66), and (67) represents an alkyl group containing a residue having any one of a phenyl structure, a thienyl structure, and a cyclohexyl structure at a terminal thereof, the substituent is more preferably selected from substituents represented by the chemical formulae (14), (15), (16), and (17).

—(CH_2) k_{14} — CH_3

(In the formula, k_{14} represents an integer selected from 0 to 8. When multiple units exist, k_{14} 's each independently have the above meaning for each unit.)

$$--(CH2)k15 --- CH3 CH3 (15)$$

(In the formula, k_{15} represents an integer selected from 0 to 7. When multiple units exist, k_{15} 's each independently have the above meaning for each unit.) $-(CH_2)k_{16}-R_{16}$ (16)

(In the formula, k_{16} represents an integer selected from 1 to 8. R_{16} represents a substituent containing a residue having any one of a phenyl structure and a thienyl structure. When multiple units exist, k_{16} and R_{16} each independently have the above meaning for each unit.)

$$-(CH_2)k_{17}$$

(In the formula, R_{17} represents a substituent to a cyclohexyl group selected from an H atom, a CN group,

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an NO₂ group, a halogen atom, a CH₃ group, a C_2H_5 group, a C_3H_7 group, a CF₃ group, a C_2F_5 group, and a C_3F_7 group. k_{17} represents an integer selected from 0 to 8. When multiple units exist, k_{17} and R_{17} each independently have the above meaning for each unit.)

In addition, R_{16} in the chemical formula (16), that is, a residue having any one of a phenyl structure and a thienyl structure is selected from the group of residues represented by the chemical formulae (18), (19), (20), (21), (22), (23), (24), (25), (26), (27), and (28).

Here, the chemical formula (18) represents a group of unsubstituted or substituted phenyl groups.

(In the formula, R₁₈ represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO₂ group, a CH₃ group, a C₂H₅ group, a C₃H₇ group, a CH=CH₂ group, COOR_{18a} (R_{18a} represents an H atom, an Na atom, or a K atom.), a CF₃ group, a C₂F₅ group, and a C₃F₇ group. When multiple units exist, R₁₈'s may be different for each unit.)

The chemical formula (19) represents a group of unsubstituted or substituted phenoxy groups.

(In the formula, R_{19} represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a SCH_3 group, a CF_3 group, a C_2F_5 group, and a C_3F_7 group. When multiple units exist, R_{19} 's may be different for each unit.)

The chemical formula (20) represents a group of unsubstituted or substituted benzoyl groups.

(In the formula, R_{20} represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a CF_3 group, a C_2F_5 group, and a C_3F_7 group.

15 When multiple units exist, R_{20} 's may be different for each unit.)

The chemical formula (21) represents a group of unsubstituted or substituted phenylsulfanyl groups.

20 (In the formula, R_{21} represents a substituent to an aromatic ring selected from an H atom, a halogen atom,

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a CN group, an NO₂ group, COOR_{21a}, SO₂R_{21b} (R_{21a} represents H, Na, K, CH₃, or C₂H₅, and R_{21b} represents OH, ONa, OK, a halogen atom, OCH₃, or OC₂H₅.), a CH₃ group, a C₂H₅ group, a C₃H₇ group, a (CH₃)₂-CH group, and a (CH₃)₃-C group. When multiple units exist, R₂₁'s may be different for each unit.)

The chemical formula (22) represents a group of unsubstituted or substituted (phenylmethyl)sulfanyl groups.

$$R_{22}$$
 $-CH_2$ $-S$ (22)

(In the formula, R₂₂ represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO₂ group, COOR_{22a}, SO₂R_{22b} (R_{22a} represents H, Na, K, CH₃, or C₂H₅, and R_{22b} represents

OH, ONa, OK, a halogen atom, OCH₃, or OC₂H₅.), a CH₃ group, a C₂H₅ group, a C₃H₇ group, a (CH₃)₂-CH group, and a (CH₃)₃-C group. When multiple units exist, R₂₂'s may be different for each unit.)

The chemical formula (23) represents a 2-20 thienyl group.

The chemical formula (24) represents a 2-thienylsulfanyl group.

The chemical formula (25) represents a 2-thienylcarbonyl group.

5 The chemical formula (26) represents a group of unsubstituted or substituted phenylsulfinyl groups.

(In the formula, R₂₆ represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO₂ group, COOR_{26a}, SO₂R_{26b} (R_{26a} represents H, Na, K, CH₃, or C₂H₅, and R_{26b} represents OH, ONa, OK, a halogen atom, OCH₃, or OC₂H₅.), a CH₃ group, a C₂H₅ group, a C₃H₇ group, a (CH₃)₂-CH group, and a (CH₃)₃-C group. When multiple units exist, R₂₆'s may be different for each unit.)

The chemical formula (27) represents a group of unsubstituted or substituted phenylsulfonyl groups.

(In the formula, R_{27} represents a substituent to 20 an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO_2 group, $COOR_{27a}$, SO_2R_{27b} (R_{27a}

represents H, Na, K, CH_3 , or C_2H_5 , and R_{27b} represents OH, ONa, OK, a halogen atom, OCH₃, or OC₂H₅.), a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $(CH_3)_2$ -CH group, and a $(CH_3)_3$ -C group. When multiple units exist, R_{27} 's may be different for each unit.)

The chemical formula (28) represents a (phenylmethyl)oxy group.

Examples of the compound represented by the chemical formula (34) include methyl chloroformate, 10 ethyl chloroformate, propyl chloroformate, isopropyl chloroformate, butyl chloroformate, cyclohexyl chloroformate, benzyl chloroformate, methyl bromoformate, ethyl bromoformate, propyl bromoformate, 15 isopropyl bromoformate, butyl bromoformate, cyclohexyl bromoformate, benzyl bromoformate, methyl chloroacetate, ethyl chloroacetate, propyl chloroacetate, isopropyl chloroacetate, butyl chloroacetate, cyclohexyl chloroacetate, benzyl chloroacetate, methyl bromoacetate, ethyl 20 bromoacetate, propyl bromoacetate, isopropyl bromoacetate, butyl bromoacetate, cyclohexyl bromoacetate, benzyl bromoacetate, methyl 3chloropropionate, ethyl 3-chloropropionate, propyl 3-25 chloropropionate, isopropyl 3-chloropropionate, butyl

3-chloropropionate, cyclohexyl 3-chloropropionate, benzyl 3-chloropropionate, methyl 3-bromopropionate, ethyl 3-bromopropionate, propyl 3-bromopropionate, isopropyl 3-bromopropionate, butyl 3-bromopropionate, cyclohexyl 3-bromopropionate, benzyl 3-5 bromopropionate, methyl 4-chlorobutyrate, ethyl 4chlorobutyrate, propyl 4-chlorobutyrate, isopropyl 4chlorobutyrate, butyl 4-chlorobutyrate, cyclohexyl 4chlorobutyrate, benzyl 4-chlorobutyrate, methyl 4-10 bromobutyrate, ethyl 4-bromobutyrate, propyl 4bromobutyrate, isopropyl 4-bromobutyrate, butyl 4bromobutyrate, cyclohexyl 4-bromobutyrate, benzyl 4bromobutyrate, methyl 5-chlorovalerate, ethyl 5chlorovalerate, propyl 5-chlorovalerate, isopropyl 5-15 chlorovalerate, butyl 5-chlorovalerate, cyclohexyl 5chlorovalerate, benzyl 5-chlorovalerate, methyl 5bromovalerate, ethyl 5-bromovalerate, propyl 5bromovalerate, isopropyl 5-bromovalerate, butyl 5bromovalerate, cyclohexyl 5-bromovalerate, benzyl 5-20 bromovalerate, methyl 6-chlorohexanoate, ethyl 6chlorohexanoate, propyl 6-chlorohexanoate, isopropyl 6-chlorohexanoate, butyl 6-chlorohexanoate, cyclohexyl 6-chlorohexanoate, benzyl 6chlorohexanoate, methyl 6-bromohexanoate, ethyl 6-25 bromohexanoate, propyl 6-bromohexanoate, isopropyl 6bromohexanoate, butyl 6-bromohexanoate, cyclohexyl 6bromohexanoate, benzyl 6-bromohexanoate, methyl 7-

chloroheptanoate, ethyl 7-chloroheptanoate, propyl 7chloroheptanoate, isopropyl 7-chloroheptanoate, butyl 7-chloroheptanoate, cyclohexyl 7-chloroheptanoate, benzyl 7-chloroheptanoate, methyl 7-bromoheptanoate, ethyl 7-bromoheptanoate, propyl 7-bromoheptanotate, isopropyl 7-bromoheptanoate, butyl 7-bromoheptanoate, cyclohexyl 7-bromoheptanoate, benzyl 7-bromooctanoate, methyl 8-chlorooctanoate, ethyl 8-chlorooctanoate, propyl 8-chlorooctanoate, isopropyl 8-chlorooctanoate, butyl 8-chlorooctanotate, cyclohexyl 8-10 chlorooctanoate, benzyl 8-chlorooctanoate, methyl 8bromooctanoate, ethyl 8-bromooctanoate, propyl 8bromooctanoate, isopropyl 8-bromooctanoate, butyl 8bromooctanoate, cyclohexyl 8-bromooctanoate, benzyl 8-bromooctanoate, methyl 9-chlorononanoate, ethyl 9-15 chlorononanoate, propyl 9-chlorononanoate, isopropyl 9-chlorononanoate, butyl 9-bromononanoate, cyclohexyl 9-chlorononanoate, benzyl 9-chlorononanoate, methyl 9-bromononanoate, ethyl 9-bromononanoate, propyl 9bromononanoate, isopropyl 9-bromononanoate, butyl 9-20 bromononanoate, cyclohexyl 9-bromononanoate, and benzyl 9-bromononanoate.

Method of producing polyhydroxyalkanoate represented by chemical formula (35)

A reaction between the polyhydroxyalkanoate containing a unit represented by the chemical formula (33) and the compound represented by the chemical

formula (34) in the present invention will be described in detail.

The present invention can be achieved by subjecting a α -methylene or a α -methine adjacent to a carbonyl group in a polymer main chain to an 5 addition reaction with the compound represented by the chemical formula (34). To be specific, the present invention can be achieved by: allowing the polyhydroxyalkanoate containing a unit represented by the chemical formula (33) to react with a base 10 capable of forming a α -methylene or a α -methine, which is adjacent to a carbonyl group in the polymer main chain of the polyhydroxyalkanoate containing a unit represented by the chemical formula (33), into 15 an anion under an addition reaction condition; and allowing the resultant to react with the compound represented by the chemical formula (34). In the present invention, the amount of the compound represented by the chemical formula (34) to be used is 0.001 to 100 times mole, or preferably 0.01 to 10 20 times mole with respect to the unit represented by the chemical formula (33).

A solvent to be used in the reaction of the present invention is not particularly limited as long as it is inactive to the reaction and dissolves the staring material to some extent. Examples of such a solvent include: aliphatic hydrocarbons such as

hexane, cyclohexane, heptane, ligroin, and petroleum ether; aromatic hydrocarbons such as benzene, toluene, and xylene; ethers such as diethyl ether, diisopropyl ether, tetrahydrofuran, dioxane, dimethoxyethane, and diethyleneglycoldimethylether; and amides such as formamide, N,N-dimethylformamide, N,N-dimethylacetamide, N-methyl-2-pyrrolidone, N-methylpyrrolidinone, and hexamethylphosphorotriamide. Of those, tetrahydrofuran is preferable.

The reaction is performed in the presence of a 10 base. Examples of a base to be used include: lithium alkyls such as methyl lithium and butyl lithium; alkali metal disilazides such as lithium hexamethyl disilazide, sodium hexamethyl disilazide, and potassium hexamethyl disilazide; and lithium amides 15 such as lithium diisopropylamide and lithium dicyclohexylamide. Of those, lithium diisopropylamide is preferable. In addition, the amount of the base to be used in the present invention is 0.001 to 100 times mole, or preferably 20 0.01 to 10 times mole with respect to the unit represented by the chemical formula (33).

In the method of the present invention, a reaction temperature is generally in the range of -78°C to 40°C, or preferably in the range of -78°C to 30°C.

In the method of the present invention, a

reaction time is generally in the range of 10 minutes to 24 hours. The reaction time is particularly preferably in the range of 10 minutes to 4 hours.

The polyhydroxyalkanoate having a unit represented by the chemical formula (33) included in the chemical formula (5) can be produced according to the above production method.

In addition, a polymer produced by means of a conventionally known method can be arbitrarily used as the polyhydroxyalkanoate containing a unit 10 represented by one of the chemical formulae (33) and (36) to be used in the present invention. Examples of a polyhydroxyalkanoate represented by the chemical formula (68) included in the chemical formulae (33) and (35) include organism-produced polyesters 15 typified by poly-3-hydroxybutyrate (k68 in the chemical formula (68) represents 0), poly-3hydroxyvalerate (k68 in the chemical formula (68) represents 1), and the like. For example, JP-B H07-14352 and JP-B H08-19227 each disclose a method of 20 producing a copolymer of 3-hydroxybutyrate and 3hydroxyvalerate. In addition, JP-A H05-93049 and JP-A H07-265065 each disclose a method of producing a copolymer of 3-hydroxybutyrate and 3-hydroxyhexanoate (k_{68} represents 2). In addition, JP 2642937 B 25 discloses a method of producing a copolymer containing a 3-hydroxyalkanoate having 6 to 12 carbon

atoms (that is, from 3-hydroxyhexanoate to 3hydorxyundecanoate). JP-A 2002-306190 discloses a method of producing a homopolymer of poly-3hydroxybutyrate. A polyhydroxyalkanoate can be produced in the present invention by means of a In addition, a polyhydroxyalkanoate similar method. containing a unit represented by the chemical formula (69) included in the chemical formulae (33) and (35) can be produced by means of a method disclosed in 10 International Journal of Biological Macromolecules 12 (1990) 92. In addition, a method of producing a polyhydroxyalkanoate containing a unit represented by the chemical formula (70) or (71) included in the chemical formulae (33) and (35) is disclosed in JP-A 15 2001-288256 and JP-A 2003-319792. A polyhydroxyalkanoate can be produced in the present invention by means of a similar method.

$$\begin{array}{c|c}
 & H & H \\
\hline
 & H & CH_2 \\
 & CH_3
\end{array}$$
(68)

 $(k_{68} \text{ represents an integer, selected from 0 to 8.}$ 20 When multiple units exist, k_{68} 's each independently have the above meaning for each unit.)

15

$$\begin{array}{c|c}
 & H & H \\
\hline
O & H_{(CH_2)k_{69}} \\
 & CH \\
 & H_3C & CH_3
\end{array}$$
(69)

 $(k_{69} \text{ represents an integer selected from 0 to 7.}$ When multiple units exist, k_{69} 's each independently have the above meaning for each unit.)

$$\begin{array}{c|c}
 & H & H \\
\hline
 & H & C \\
 & H & C \\
 & C & C \\
 & R_{70}
\end{array}$$
(70)

 $(k_{70} \ \text{represents} \ \text{an integer selected from 1 to 8.} \ R_{70}$ represents a substituent containing a residue having any one of a phenyl structure and a thienyl structure. When multiple units exist, k_{70} and R_{70} each

10 independently have the above meaning for each unit.)

$$\begin{array}{c|c}
H & H \\
\hline
O & H (CH_2)k_{71}
\end{array}$$

$$\begin{array}{c|c}
R_{71}
\end{array}$$
(71)

(In the formula, R_{71} represents a substituent to a cyclohexyl group selected from an H atom, a CN group, an NO_2 group, a halogen atom, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a CF_3 group, a C_2F_5 group, and a C_3F_7 group. k_{71} represents an integer selected from O

15

to 8. When multiple units exist, k_{71} and R_{71} each independently have the above meaning for each unit.)

In addition, R_{70} in the chemical formula (70), that is, a residue having any one of a phenyl structure and a thienyl structure is selected from the group of residues represented by the chemical formulae (72), (73), (74), (75), (76), (77), (78), (79), (80), (81), and (82).

Here, the chemical formula (72) represents a group of unsubstituted or substituted phenyl groups.

(In the formula, R_{72} represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a $CH=CH_2$ group, $COOR_{72a}$ (R_{72a} represents an H atom, an Na atom, or a K atom.), a CF_3 group, a C_2F_5 group, and a C_3F_7 group. When multiple units exist, R_{72} 's may be different for each unit.)

The chemical formula (73) represents a group of unsubstituted or substituted phenoxy groups.

(In the formula, R_{73} represents a substituent to an aromatic ring selected from an H atom, a halogen atom,

10

20

a CN group, an NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a SCH_3 group, a CF_3 group, a C_2F_5 group, and a C_3F_7 group. When multiple units exist, R_{73} 's may be different for each unit.)

The chemical formula (74) represents a group of unsubstituted or substituted benzoyl groups.

(In the formula, R_{74} represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO_2 group, a CH_3 group, a C_2H_5 group, a C_3H_7 group, a CF_3 group, a C_2F_5 group, and a C_3F_7 group. When multiple units exist, R_{74} 's may be different for each unit.)

The chemical formula (75) represents a group of unsubstituted or substituted phenylsulfanyl groups.

(In the formula, R_{75} represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an_NO₂_group,_COOR_{75a}, SO₂R_{75b} (R_{75a} represents H, Na, K, CH₃, or C₂H₅, and R_{75b} represents OH, ONa, OK, a halogen atom, OCH₃, or OC₂H₅.), a CH₃ group, a C₂H₅ group, a C₃H₇ group, a (CH₃)₂-CH group, and a (CH₃)₃-C group. When multiple units exist, R₇₅'s

may be different for each unit.)

The chemical formula (76) represents a group of unsubstituted or substituted (phenylmethyl)sulfanyl groups.

$$R_{76}$$
 CH_2 $-S$ (76)

(In the formula, R₇₆ represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO₂ group, COOR_{76a}, SO₂R_{76b} (R_{76a} represents H, Na, K, CH₃, or C₂H₅, and R_{76b} represents

OH, ONa, OK, a halogen atom, OCH₃, or OC₂H₅.), a CH₃ group, a C₂H₅ group, a C₃H₇ group, a (CH₃)₂-CH group, and a (CH₃)₃-C group. When multiple units exist, R₇₆'s may be different for each unit.)

The chemical formula (77) represents a 2-15 thienyl group.

The chemical formula (78) represents a 2-thienylsulfanyl group.

20 The chemical formula (79) represents a 2-thienylcarbonyl group.

The chemical formula (80) represents a group of unsubstituted or substituted phenylsulfinyl groups.

(In the formula, R₈₀ represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO₂ group, COOR_{80a}, SO₂R_{80b} (R_{80a} represents H, Na, K, CH₃, or C₂H₅, and R_{80b} represents OH, ONa, OK, a halogen atom, OCH₃, or OC₂H₅.), a CH₃ group, a C₂H₅ group, a C₃H₇ group, a (CH₃)₂-CH group, and a (CH₃)₃-C group. When multiple units exist, R₈₀'s may be different for each unit.)

The chemical formula (81) represents a group of unsubstituted or substituted phenylsulfonyl groups.

(In the formula, R₈₁ represents a substituent to an aromatic ring selected from an H atom, a halogen atom, a CN group, an NO₂ group, COOR_{81a}, SO₂R_{81b} (R_{81a} represents H, Na, K, CH₃, or C₂H₅, and R_{81b} represents OH, ONa, OK, a halogen atom, OCH₃, or OC₂H₅.), a CH₃ group, a C₂H₅ group, a C₃H₇ group, a (CH₃)₂-CH group, and a (CH₃)₃-C group. When multiple units exist, R₈₁'s may be different for each unit.)

The chemical formula (82) represents a group of

unsubstituted or substituted (phenylmethyl)oxy groups.

(82)

The novel polyhydroxyalkanoate and the method of producing the same shown in the present invention are provided by using a polyhydroxyalkanoate 5 containing a unit represented by the chemical formula (33) or (35) including the chemical formulae (68), (69), (70), and (71) as a starting material. However, the present invention is not limited to the method described above. 10

The molecular weight of the polyhydroxyalkanoate of the present invention can be measured as a relative molecular weight or an absolute molecular weight. The molecular weight can 15 be simply measured by means of, for example, gel permeation chromatography (GPC). A specific measurement method by means of GPC is as follows. The polyhydroxyalkanoate is dissolved in advance into a solvent into which the polyhydroxyalkanoate is soluble, and the molecular weight is measured in a 20 mobile phase of the same solvent. A differential refractometer (RI) or an ultraviolet (UV) detector can be used as a detector depending on the polyhydroxyalkanoate to be measured. The molecular weight is determined as a result of relative 25

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comparison with a standard sample (such as polystyrene or polymethyl methacrylate). The solvent can be selected from solvents into each of which a polymer is soluble such as dimethylformamide (DMF), dimethyl sulfoxide (DMSO), chloroform, tetrahydrofuran (THF), toluene, and hexafluoroisopropanol (HFIP). In the case of a polar solvent, the molecular weight can be measured through addition of a salt.

The number average molecular weight of a 10 polyhydroxyalkanoate to be produced by the present invention can vary widely by changing conditions including a reaction time, a reaction temperature, and a reaction time. The optimum number average molecular weight of the polyhydroxyalkanoate, which 15 varies depending on a target function, is in the range of 1,000 to 1,000,000 when one attempts to use the polyhydroxyalkanoate for a medical soft member or the like. In addition, a polyhydroxyalkanoate having a ratio (Mw/Mn) between a weight average molecular 20 weight (Mw) and the number average molecular weight (Mn) in the range of 1 to 10 is preferable.

A reaction solvent, a reaction temperature, a reaction time, a purification method, and the like in a chemical reaction of the present invention are not limited to those described above.

25

[EXAMPLES]

Hereinafter, the present invention will be described in more detail by way of examples. However, the present invention is not limited to these examples.

In each of Examples 1 to 4, a microorganism is used to produce a polyhydroxyalkanoate. The microorganisms used in those examples are a Ralstonia eutropha TB64 strain (disclosed in JP-A 2000-166587) and a Pseudomonas cichorii YN2 strain (FERM BP-7375, disclosed in JP-A 2001-288256). Those 2 microorganisms are deposited in the National Institute of Advanced Industrial Science and Technology, International Patent Organism Depositary.

The mineral salt medium (M9 medium) used in each of Examples 1 to 4 has the following composition.

M9 medium composition (in 1 L)

 Na_2HPO_4 6.2 g KH_2PO_4 3.0 g

20 NaCl 0.5 g NH_4Cl 1.0 g

Water Balance

(pH 7.0)

For better proliferation of a microorganism and better production of a polyhydroxyalkanoate at the time of culture, the above mineral salt is added with about 0.3% (volume/volume) of a trace component

solution shown below.

(Trace component solution composition: unit g/L)
Nitrilotriacetic acid: 1.5; MgSO₄: 3.0; MnSO₄: 0.5;
NaCl: 1.0; FeSO₄: 0.1; CaCl₂: 0.1; CoCl₂: 0.1; ZnSO₄:
0.1; CuSO₄: 0.1; AlK(SO₄)₂: 0.1; H₃BO₃: 0.1; Na₂MoO₄:
0.1; NiCl₂: 0.1

[Example 1]

(Synthesis of poly-3-hydroxybutyric acid represented by chemical formula (101))

Poly-3-hydroxybutyric acid represented by the chemical formula (101) was synthesized by means of the method disclosed in Example 1 of JP-A 2002-306190.

A colony of a TB 64 strain on an M9 agar medium containing 0.1% of sodium malate was inoculated in 50 ml of an M9 medium containing 0.5% of sodium malate in a 500-mL shaking flask, and the whole was shake cultured at 30°C. 24 hours after that, 5 ml of the culture solution were added to 1 L of a production medium prepared by incorporating 0.5% of sodium malate into an M9 medium with the concentration of only NH₄Cl as a nitrogen source reduced to 1/10, and the whole was shaken in the same manner to accumulate PHB in the cells. 48 hours after that, the cells

were recovered by centrifugal separation, washed with methanol, and then freeze-dried. After the dried cells had been weighed, chloroform was added, and the whole was stirred at 60°C for 24 hours to extract a polymer.

After filtrating the extracted chloroform solution through a filter, it was concentrated by means of an evaporator. After that, a portion precipitated and solidified with cold methanol was collected and dried under reduced pressure to prepare 1.83 g of a polymer per L of the production medium. NMR analysis was performed under the following conditions to determine the structure of the resultant polymer.

<Measuring equipment> FT-NMR: Bruker DPX 400

15 Resonance frequency: ¹H = 400 MHz

Measurement conditions> Measured nuclear species: ¹H
Solvent used: CDCl₃

Measurement temperature: room temperature

The analysis confirmed that the resultant

20 polymer was a polyhydroxyalkanoate containing a unit
of 3-hydroxybutyric acid represented by the chemical
formula (1). The resultant polyhydroxyalkanoate was
evaluated for average molecular weight by means of
gel permeation chromatography (GPC; HLC-8220

25 manufactured by Tosoh Corporation, column; TSK-GEL
Super HM-H manufactured by Tosoh Corporation,

solvent; chloroform, polystyrene conversion). As a

10

result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 549,500 and a weight average molecular weight Mw of 1,263,900.

45.6 g of the polyhydroxyalkanoate to be used in each of Examples 5 to 8 was prepared from 50 L of the production medium by means of the above method.

[Example 2]

(Synthesis of poly-3-hydroxy-5-phenylvaleric acid represented by chemical formula (102))

Poly-3-hydroxy-5-phenylvaleric acid represented by the chemical formula (102) was synthesized by means of the method disclosed in Example 1 of JP-A 2003-319792. 200 mL of an M9 medium containing 0.5% (weight/volume (w/v)) of polypeptone (Wako Pure Chemical Industries, Ltd.) and 0.1% (weight/volume (w/v)) of 5-phenylvaleric acid were prepared as a production medium. 1 mL of a culture solution prepared in advance by shake culturing a Pseudomonas cichorii YN2 strain in an M9 medium containing 0.5% of polypeptone at 30°C for 8 hours was added to the

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production medium, and the whole was cultured in a 500-mL shaking flask at 30°C for 24 hours. After the culture, the cells were recovered by centrifugal separation, washed with methanol, and then freezedried. After the dried cells had been weighed, 5 chloroform was added, and the whole was stirred at 50°C for 24 hours to extract a polymer. After filtrating the extracted chloroform solution through a filter, it was concentrated by means of an evaporator. After that, a portion precipitated and 10 solidified with cold methanol was collected and dried under reduced pressure to prepare 0.60 g of a polymer per L of the production medium. NMR analysis was performed under the same conditions as those of Example 1 to determine the structure of the resultant 15 polymer. The analysis confirmed that the resultant polymer was substantially a homopolymer of a unit of poly-3-hydroxy-5-phenylvaleric acid represented by the chemical formula (102) as a monomer unit. The average molecular weight of the resultant 20 polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 91,000 and a weight average molecular weight Mw of 172,900. 25

60.1 q of the polyhydroxyalkanoate to be used in each of Examples 9 to 12 was prepared from 100 L

of the production medium by means of the above method. [Example 3]

(Synthesis of poly-3-hydroxy-5-phenoxyvaleric acid represented by chemical formula (103))

Poly-3-hydroxy-5-phenylvaleric acid represented by the chemical formula (103) was synthesized by means of the method disclosed in Example 4 of JP-A 2003-319792.

(weight/volume (w/v)) of polypeptone (Wako Pure Chemical Industries, Ltd.) and 0.1% (weight/volume (w/v)) of 5-phenoxyvaleric acid was prepared as a production medium. 1 mL of a culture solution

15 prepared in advance by shake culturing a Pseudomonas cichorii YN2 strain in an M9 medium containing 0.5% of polypeptone at 30°C for 8 hours was added to the production medium, and the whole was cultured in a 500-mL shaking flask at 30°C for 45 hours. After the culture, the cells were recovered by centrifugal separation, washed with methanol, and then freeze-

dried. After the dried cells had been weighed, chloroform was added, and the whole was stirred at 50°C for 24 hours to extract a polymer. After filtrating the extracted chloroform solution through a filter, it was concentrated by means of an 5 evaporator. After that, a portion precipitated and solidified with cold methanol was collected and dried under reduced pressure to prepare 0.36 g of a polymer per L of the production medium. NMR analysis was performed under the same conditions as those of 10 Example 1 to determine the structure of the resultant polymer. The analysis confirmed that the resultant polymer was substantially a homopolymer of a unit of poly-3-hydroxy-5-phenoxyvaleric acid represented by the chemical formula (103) as a monomer unit. The 15 average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 201,000 and a 20 weight average molecular weight Mw of 422,100.

 $44.8\ g$ of the polyhydroxyalkanoate to be used in each of Examples 13 to 16 was prepared from 125 L of the production medium by means of the above method.

25 [Example 4]

(Synthesis of poly-3-hydroxy-4-cyclohexylbutyric acid represented by chemical formula (104)

Poly-3-hydroxy-4-cyclohexylbutyric acid represented by the chemical formula (104) was synthesized by means of the method disclosed in Example 9 of JP-A 2003-319792.

200 mL of an M9 medium containing 0.5% (weight/volume (w/v)) of polypeptone (Wako Pure Chemical Industries, Ltd.) and 0.1% (weight/volume (w/v)) of 4-cyclohexylbutyric acid was prepared as a production medium. 1 mL of a culture solution 10 prepared in advance by shake culturing a Pseudomonas cichorii YN2 strain in an M9 medium containing 0.5% of polypeptone at 30°C for 8 hours was added to the production medium, and the whole was cultured in a 500-mL shaking flask at 30°C for 48 hours. After the 15 culture, the cells were recovered by centrifugal separation, washed with methanol, and then freezedried. After the dried cells had been weighed, chloroform was_added_r -and_the whole was stirred at 50°C for 24 hours to extract a polymer. After 20 filtrating the extracted chloroform solution through a filter, it was concentrated by means of an evaporator. After that, a portion precipitated and

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solidified with cold methanol was collected and dried under reduced pressure to prepare 0.48 g of a polymer per L of the production medium. NMR analysis was performed under the same conditions as those of Example 1 to determine the structure of the resultant 5 polymer. The analysis confirmed that the resultant polymer was substantially a homopolymer of a unit of poly-3-hydroxy-4-cyclohexylbutyric acid represented by the chemical formula (104) as a monomer unit. The average molecular weight of the resultant 10 polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 70,500 and a weight average molecular weight Mw of 155,100. 15

47.9 g of the polyhydroxyalkanoate to be used in each of Examples 17 and 18 was prepared from 100 L of the production medium by means of the above method. [Example 5]

10.00 g of the polyhydroxyalkanoate composed of the unit represented by the chemical formula (101) synthesized in Example 1 was placed in a round-bottomed flask, and 500 ml of THF was added to dissolve this. The flask was placed under a nitrogen atmosphere, and the solution was stirred at -78°C.

Next, 58.08 ml (116.2 mmol) of a solution of 2 M of lithium diisopropylamide in THF was gradually added

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to the flask, and the whole was stirred at $-78\,^{\circ}\text{C}$ for 30 minutes. Next, 19.82 g (232.3 mmol) of benzyl chloroformate was added to the flask, and the whole was stirred at room temperature for 30 minutes.

- 5 After the completion of the reaction, the reaction solution was poured into 1,000 ml of an aqueous solution of ammonium chloride, and 500 ml of dichloromethane was added to extract the organic layer. The extracted organic layer was washed with
- 250 ml of water 3 times. After the organic layer had been collected, the solvent was distilled off to collect a crude polymer. Next, the polymer was dissolved into 60 ml of THF, and reprecipitated in methanol in an amount 50 times that of THF necessary
- 15 for the dissolution. The precipitate was collected and dried under reduced pressure to prepare 8.44 g of a polymer. NMR analysis was performed under the same conditions as those of Example 1 to determine the structure of the resultant polymer. The analysis
- confirmed that the polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (105) as a monomer unit. The analysis also confirmed that an A unit accounted for 10 mol% of the monomer unit and a B unit accounted

25

for 90 mol% thereof.

The resultant polyhydroxyalkanoate was
evaluated for average molecular weight by means of
gel permeation chromatography (GPC; HLC-8220

5 manufactured by Tosoh Corporation, column; TSK-GEL
Super HM-H manufactured by Tosoh Corporation,
solvent; chloroform, polystyrene conversion). As a
result, the resultant polyhydroxyalkanoate was found
to have a number average molecular weight Mn of
325,400 and a weight average molecular weight Mw of
764,700.

5.00 g of the polyhydroxyalkanoate copolymer represented by the chemical formula (105) synthesized here was dissolved into 500 ml of a mixed solvent of dioxane-ethanol (75 : 25), and 1.10 g of a 5% palladium/carbon catalyst was added to the solution. The inside of the reaction system was filled with hydrogen, and the whole was stirred at room temperature for 1 day. After the completion of the reaction, in order to remove the catalyst, the resultant was filtered through a 0.25-µm membrane

15

filter to collect a reaction solution. After the solution had been concentrated, the concentrate was dissolved into chloroform, and reprecipitated in methanol in an amount 10 times that of chloroform. The resultant polymer was collected and dried under reduced pressure to prepare 3.59 g of a polymer. NMF analysis was performed under the same conditions as those of Example 6 to determine the structure of the resultant polymer. The analysis confirmed that the polymer was a polyhydroxyalkanoate copolymer containing a unit represented by the following chemical formula (106) as a monomer unit. The analysis also confirmed that a C unit accounted for

10 mol% of the monomer unit and a D unit accounted

for 90 mol% thereof.

The resultant polyhydroxyalkanoate was
evaluated for average molecular weight by means of
gel permeation chromatography (GPC; HLC-8220

20 manufactured by Tosoh Corporation, column; TSK-GEL
Super HM-H manufactured by Tosoh Corporation,
solvent; chloroform, polystyrene conversion). As a
result, the resultant polyhydroxyalkanoate was found
to have a number average molecular weight Mn of

298,000 and a weight average molecular weight Mw of 715,200.

Furthermore, 30 mg of the polyhydroxyalkanoate synthesized here was placed in a 100-ml roundbottomed flask, and 2.1 ml of chloroform and 0.7 ml of methanol were added to dissolve this. 0.5 ml of a 2-mol/L trimethylsilyldiazomethane-hexane solution was added to the solution, and the whole was stirred at room temperature for 1 hour. After the completion of the reaction, the solvent was distilled off to 10 collect a polymer. The polymer was washed with 50 ml of methanol to collect a polymer. The polymer was dried under reduced pressure to prepare 29 mg of a polyhydroxyalkanoate. The resultant polyhydroxyalkanoate was subjected to NMR analysis in 15 the same manner as in Example 1. The analysis confirmed that a carboxyl group of the C unit was transformed into methyl carboxylate, and that the resultant polymer can be esterified again:

20 [Example 6]

25

9.40 g of a polymer was prepared in the same manner as in Example 5 except that 26.61 g (232.3 mmol) of benzyl bromoacetate was used instead of benzyl chloroformate. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 1. The analysis confirmed that the polymer was a polyhydroxyalkanoate containing a

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15

20

unit represented by the following chemical formula (107). The analysis also confirmed that an A unit accounted for 11 mol% of the monomer unit and a B unit accounted for 89 mol% thereof.

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 300,300 and a weight average molecular weight Mw of 723,700.

The above polymer was subjected to hydrogenolysis in the same manner as in Example 5 to prepare 3.66 g of a polymer. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 1. The analysis confirmed that the polymer was a polyhydroxyalkanoate copolymer containing a unit represented by the following chemical formula (108) as a monomer unit. The analysis also confirmed that a C unit accounted

15

20

for 11 mol% of the monomer unit and a D unit accounted for 89 mol% thereof.

$$\begin{array}{c|c}
COOH \\
CH_2H \\
O H CH_3
\end{array}$$

$$\begin{array}{c|c}
O H H \\
C D
\end{array}$$

$$\begin{array}{c|c}
O H H \\
CH_3
\end{array}$$

$$\begin{array}{c|c}
O H CH_3
\end{array}$$

$$\begin{array}{c|c}
O H CH_3
\end{array}$$

$$\begin{array}{c|c}
O H CH_3
\end{array}$$

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 286,000 and a weight average molecular weight Mw of 700,700.

10 [Example 7]

manner as in Example 5 except that 22.66 g (232.3 mmol) of ethyl 4-bromobutylate was used instead of benzyl chloroformate. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 1. The analysis confirmed that the polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (109). The analysis also confirmed that an A unit accounted for 10 mol% of the monomer unit and a B unit accounted for 90 mol% thereof.

10

15

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 300,300 and a weight average molecular weight Mw of 723,700.

The above polymer was subjected to hydrogenolysis in the same manner as in Example 5 to prepare 3.93 g of a polymer. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 1. The analysis confirmed that the polymer was a polyhydroxyalkanoate copolymer containing a unit represented by the following chemical formula (110) as a monomer unit. The analysis also confirmed that a C unit accounted for 10 mol% of the monomer unit and a D unit accounted for 90 mol% thereof.

$$\begin{array}{c|c}
COOH \\
(CH_2)_{3H} \\
\hline
(CH_2)_{3H} \\
O H CH_3
\end{array}$$

$$\begin{array}{c|c}
O H H \\
H CH_3
\end{array}$$

$$\begin{array}{c|c}
O \\
O H CH_3
\end{array}$$

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 286,000 and a weight average molecular weight Mw of 700,700. [Example 8]

8.83 g of a polymer was prepared in the same
10 manner as in Example 5 except that 29.17 g (232.3 mmol) of ethyl 8-bromooctanoate was used instead of benzyl chloroformate. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 1. The analysis confirmed that the polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (111). The analysis also confirmed that an A unit accounted for 9 mol% of the monomer unit and a B unit accounted for 91 mol% thereof.

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 321,000 and a weight average molecular weight Mw of 776,800.

The above polymer was subjected to
hydrogenolysis in the same manner as in Example 5 to
10 prepare 3.85 g of a polymer. The resultant polymer
was subjected to NMR analysis under the same
conditions as those of Example 1. The analysis
confirmed that the polymer was a polyhydroxyalkanoate
copolymer containing a unit represented by the
15 following chemical formula (112) as a monomer unit.
The analysis also confirmed that a C unit accounted
for 9 mol% of the monomer unit and a D unit accounted
for 91 mol% thereof.

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 298,100 and a weight average molecular weight Mw of 715,400.

[Example 9]

8.51 g of a polymer was prepared in the same manner as in Example 5 except that 10.00 g of the 10 polyhydroxyalkanoate composed of the unit represented by the chemical formula (102) synthesized in Example 2 instead of the polyhydroxyalkanoate composed of the unit represented by the chemical formula (101), 28.38 ml (56.8 mmol) of a solution of 2 M of lithium 15 diisopropylamide in THF, and 9.68 g (113.5 mmol) of benzyl chloroformate were used. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 1. The analysis confirmed that the polymer was a polyhydroxyalkanoate 20 containing a unit represented by the following chemical formula (113). The analysis also confirmed that an A unit accounted for 12 mol% of the monomer

unit and a B unit accounted for 88 mol% thereof.

$$\begin{array}{c|c}
 & COOH \\
 & (CH_2)_{7H} \\
\hline
 & (CH_2)_{7H} \\
\hline
 & (CH_3) \\
\hline$$

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 72,500 and a weight average molecular weight Mw of 141,400.

The above polymer was subjected to

hydrogenolysis in the same manner as in Example 5 to
prepare 3.72 g of a polymer. The resultant polymer
was subjected to NMR analysis under the same
conditions as those of Example 1. The analysis
confirmed that the polymer was a polyhydroxyalkanoate
copolymer containing a unit represented by the
following chemical formula (114) as a monomer unit.
The analysis also confirmed that a C unit accounted
for 12 mol% of the monomer unit and a D unit
accounted for 89-mol% -thereof.

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 69,500 and a weight average molecular weight Mw of 139,700. [Example 10]

8.37 g of a polymer was prepared in the same
10 manner as in Example 9 except that 13.00 g (113.5 mmol) of benzyl bromoacetate was used instead of benzyl chloroformate. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 1. The analysis confirmed that the polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (115). The analysis also confirmed that an A unit accounted for 12 mol% of the monomer unit and a B unit accounted for 88 mol% thereof.

10

15

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 71,000 and a weight average molecular weight Mw of 131,400.

The above polymer was subjected to hydrogenolysis in the same manner as in Example 5 to prepare 3.87 g of a polymer. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 1. The analysis confirmed that the polymer was a polyhydroxyalkanoate copolymer containing a unit represented by the following chemical formula (116) as a monomer unit. The analysis also confirmed that a C unit accounted for 12 mol% of the monomer unit and a D unit

20

accounted for 88 mol% thereof.

$$\begin{array}{c|c}
 & COOH \\
\hline
CH_2H & O & H & H \\
O & H & (CH_2)_2 & H & (CH_2)_2
\end{array}$$

$$\begin{array}{c|c}
 & C & D & (116)
\end{array}$$

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 68,000 and a weight average molecular weight Mw of 132,600. [Example 11]

7.80 g of a polymer was prepared in the same manner as in Example 9 except that 9.48 g (113.5 mmol) of methyl 3-bromopropionate was used instead of benzyl chloroformate. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 1. The analysis confirmed that 15 the polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (117). The analysis also confirmed that an A unit accounted for 11 mol% of the monomer unit and a B unit accounted for 89 mol% thereof.

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 68,500 and a weight average molecular weight Mw of 130,800.

The above polymer was subjected to
hydrogenolysis in the same manner as in Example 5 to
10 prepare 4.01 g of a polymer. The resultant polymer
was subjected to NMR analysis under the same
conditions as those of Example 1. The analysis
confirmed that the polymer was a polyhydroxyalkanoate
copolymer containing a unit represented by the
15 following chemical formula (118) as a monomer unit.
The analysis also confirmed that a C unit accounted
for 11 mol% of the monomer unit and a D unit
accounted for 89 mol% thereof.

$$\begin{array}{c|c}
COOH \\
(CH_2)_{2H} \\
\hline
O H (CH_2)_{2}
\end{array}$$

$$\begin{array}{c|c}
O H H \\
H (CH_2)_{2}
\end{array}$$

$$\begin{array}{c|c}
C D
\end{array}$$
(118)

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 67,000 and a weight average molecular weight Mw of 127,300. [Example 12]

7.87 g of a polymer was prepared in the same
10 manner as in Example 9 except that 12.86 g (113.5 mmol) of ethyl 6-bromohexanoate was used instead of benzyl chloroformate. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 1. The analysis confirmed that the polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (119). The analysis also confirmed that an A unit accounted for 8 mol% of the monomer unit and a B unit accounted for 92 mol% thereof.

10

15

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 71,000 and a weight average molecular weight Mw of 134,900.

The above polymer was subjected to hydrogenolysis in the same manner as in Example 5 to prepare 3.95 g of a polymer. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 1. The analysis confirmed that the polymer was a polyhydroxyalkanoate copolymer containing a-unit represented by the following chemical formula (120) as a monomer unit. The analysis also confirmed that a C unit accounted for 8 mol% of the monomer unit and a D unit accounted for 92 mol% thereof.

. 2

COOH
$$(CH_2)_{5H} \longrightarrow (D)$$

$$(CH_2)_{2} \longrightarrow (D)$$

$$(CH_2)_{2} \longrightarrow (D)$$

$$(120)$$

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 68,500 and a weight average molecular weight Mw of 133,600.

[Example 13]

manner as in Example 5 except that 10.00 g of the polyhydroxyalkanoate composed of the unit represented by the chemical formula (103) synthesized in Example 3 instead of the polyhydroxyalkanoate composed of the unit represented by the chemical formula (101) in

Example 5, 26.01 ml (52.0 mmol) of a solution of 2 M of lithium diisopropylamide in THF, and 8.88 g (104.1 mmol) of benzyl chloroformate were used. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 1. The

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polyhydroxyalkanoate containing a unit represented by the following chemical formula (121). The analysis also confirmed that an A unit accounted for 11 mol% of the monomer unit and a B unit accounted for 89 mol% thereof.

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 131,500 and a weight average molecular weight Mw of 282,700.

The above polymer-was subjected to hydrogenolysis in the same manner as in Example 5 to prepare 3.75 g of a polymer. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 1. The analysis

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confirmed that the polymer was a polyhydroxyalkanoate copolymer containing a unit represented by the following chemical formula (122) as a monomer unit. The analysis also confirmed that a C unit accounted for 11 mol% of the monomer unit and a D unit accounted for 89 mol% thereof.

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 121,000 and a weight average molecular weight Mw of 260,200.

[Example 14]

7.70 g of a polymer was prepared in the same manner as in Example 13 except that 11.92 g (104.1 mmol) of benzyl bromoacetate was used instead of benzyl chloroformate. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 1. The analysis confirmed that

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the polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (123). The analysis also confirmed that an A unit accounted for 11 mol% of the monomer unit and a B unit accounted for 89 mol% thereof.

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 126,500 and a weight average molecular weight Mw of 265,700.

The above polymer was subjected to hydrogenolysis in the same manner as in Example 5 to prepare 3.86 g of a polymer. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 1. The analysis

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confirmed that the polymer was a polyhydroxyalkanoate copolymer containing a unit represented by the following chemical formula (124) as a monomer unit. The analysis also confirmed that a C unit accounted for 11 mol% of the monomer unit and a D unit accounted for 89 mol% thereof.

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 116,500 and a weight average molecular weight Mw of 256,300. [Example 15]

7.56 g of a polymer was prepared in the same manner as in Example 13 except that 10.88 g (104.1 mmol) of ethyl 5-bromovalerate was used instead of benzyl chloroformate. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 1. The analysis confirmed that

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the polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (125). The analysis also confirmed that an A unit accounted for 9 mol of the monomer unit and a B unit accounted for 91 mol% thereof.

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the 10 resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 122,000 and a weight average molecular weight Mw of 270,800. The above polymer was subjected to hydrogenolysis in the same manner as in Example 5 to prepare 3.95 g of a polymer. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 1. The analysis confirmed that the polymer

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was a polyhydroxyalkanoate copolymer containing a unit represented by the following chemical formula (126) as a monomer unit. The analysis also confirmed that a C unit accounted for 9 mol% of the monomer unit and a D unit accounted for 91 mol% thereof.

COOH

$$(CH_2)_4$$
 $(CH_2)_2$
 $(CH_2)_2$
 $(CH_2)_4$
 $(CH_2)_2$
 $(CH_2)_2$
 $(CH_2)_4$
 $(CH_2)_2$
 $(CH_2)_2$
 $(CH_2)_4$
 $(CH_2)_2$
 $(CH_2)_2$

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 116,500 and a weight average molecular weight Mw of 256,300. [Example 16]

7.60 g of a polymer was prepared in the same
15 manner as in Example 13 except that 13.07 g (104.1 mmol) of ethyl 8-bromooctanoate was used instead of benzyl chloroformate. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 1. The analysis confirmed that

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the polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (127). The analysis also confirmed that an A unit accounted for 8 mol% of the monomer unit and a B unit accounted for 92 mol% thereof.

$$\begin{array}{c}
CH_{3} \\
CH_{2} \\
O \\
O \\
H_{(CH_{2})_{2}}
\end{array}$$

$$\begin{array}{c}
O \\
H \\
H_{(CH_{2})_{2}}
\end{array}$$

$$A$$

$$\begin{array}{c}
O \\
H \\
H_{(CH_{2})_{2}}
\end{array}$$

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 134,500 and a weight average molecular weight Mw of 289,200. The above polymer was subjected to hydrogenolysis in the same manner as in Example 5 to prepare 4.01 g of a polymer. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 1. The analysis confirmed that the polymer

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was a polyhydroxyalkanoate copolymer containing a unit represented by the following chemical formula (128) as a monomer unit. The analysis also confirmed that a C unit accounted for 8 mol% of the monomer unit and a D unit accounted for 92 mol% thereof.

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The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 121,000 and a weight average molecular weight Mw of 266,200. [Example 17]

7.66 g of a polymer was prepared in the same

15 manner as in Example 6 except that 10.00 g of the
polyhydroxyalkanoate composed of the unit represented
by the chemical formula (104) synthesized in Example
4 instead of the polyhydroxyalkanoate composed of
the unit represented by the chemical formula (101) in

20 Example 5, 29.72 ml (59.4 mmol) of a solution of 2 M

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of lithium diisopropylamide in THF, and 10.14 g

(118.9 mmol) of benzyl chloroformate were used. The
resultant polymer was subjected to NMR analysis under
the same conditions as those of Example 1. The

analysis confirmed that the polymer was a
polyhydroxyalkanoate containing a unit represented by
the following chemical formula (129). The analysis
also confirmed that an A unit accounted for 10 mol%
of the monomer unit and a B unit accounted for 90

mol% thereof.

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The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 54,400 and a weight average molecular weight Mw of 11,700. The above polymer was subjected to hydrogenolysis in the same manner as in Example 5 to prepare 3.85 g of a

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polymer. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 1. The analysis confirmed that the polymer was a polyhydroxyalkanoate copolymer containing a unit represented by the following chemical formula (130) as a monomer unit. The analysis also confirmed that a C unit accounted for 10 mol% of the monomer unit and a D unit accounted for 90 mol% thereof.

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 47,500 and a weight average molecular weight Mw of 103,600.

[Example 18]

7.27 g of a polymer was prepared in the same manner as in Example 17 except that 12.43 g (118.9 mmol) of ethyl 5-bromovalerate was used instead of benzyl chloroformate. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 1. The analysis confirmed that

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the polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (131). The analysis also confirmed that an A unit accounted for 9 mol% of the monomer unit and a B unit accounted for 91 mol% thereof.

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 58,500 and a weight average molecular weight Mw of 128,700. The above polymer was subjected to hydrogenolysis in the same manner as in Example 5 to prepare 4.07 g of a polymer. The resultant-polymer was subjected to NMR analysis under the same conditions as those of Example 1. The analysis confirmed that the polymer was a polyhydroxyalkanoate copolymer containing a unit represented by the following chemical formula

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(132) as a monomer unit. The analysis also confirmed that a C unit accounted for 9 mol% of the monomer unit and a D unit accounted for 91 mol% thereof.

The average molecular weight of the resultant polyhydroxyalkanoate was measured under the same conditions as those of Example 1. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 52,100 and a weight average molecular weight Mw of 114,600. [Example 19]

Under a nitrogen atmosphere, 0.40 g of the polyhydroxyalkanoate copolymer composed of the unit represented by the chemical formula (106) synthesized in Example 5 (C: 10 mol%, D: 90 mol%) and 0.24 g (1.4 mmol) of 2-aminobenzenesulfonic acid were placed in a 100-ml three-necked flask. 15.0 ml of pyridine was added to the flask, and the mixture was stirred. After that, 0.71 ml (2.7 mmol) of triphenyl phosphite was added, and the whole was heated at 120°C for 6 hours. After the completion of the reaction, the

resultant was reprecipitated in 150 ml of ethanol, followed by collection. The resultant polymer was washed with 1N hydrochloric acid for 1 day, stirred in water for 1 day to wash the polymer, and dried under reduced pressure to prepare 0.35 g of a polymer. 5 The structure of the resultant polymer was determined through analysis according to 1H-NMR (FT-NMR: Bruker DPX 400; resonance frequency: 400 MHz; measured nuclear species: 1H; solvent used: DMSO-d6; measurement temperature: room temperature) and 10 Fourier transformation-infrared absorption (FT-IR) spectrum (Nicolet AVATAR 360FT-IR). As a result of IR measurement, a peak at 1,695 cm⁻¹ derived from a carboxylic acid reduced, and a peak derived from an amide group was newly observed at 1,658 cm⁻¹. 15

¹H-NMR confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (133) as a monomer unit because a peak derived from an aromatic ring of the 2-aminobenzenesulfonic acid structure shifted.

It was also confirmed that the polyhydroxyalkanoate was a copolymer in which an E unit accounted for 10 mol% of the unit of the polyhydroxyalkanoate represented by the chemical formula (133). The resultant polymer was evaluated for average molecular weight by means of gel permeation chromatography (GPC; Tosoh Corporation HLC-8120, column; Polymer Laboratories PLgel 5 µ MIXED-C, solvent; DMF/LiBr 0.1% (w/v), in terms of polystyrene). As a result, the resultant polymer was found to have a number average molecular weight Mn of 226,000 and a weight average molecular weight Mw of 497,200.

[Example 20]

0.33 g of a polymer was prepared in the same 15 manner as in Example 19 except that 0.28 g (1.4 mmol) of 4-methoxyaniline-2-sulfonic acid was used instead of 2-aminobenzenesulfonic acid in Example 19. The resultant polymer was subjected to NMR analysis and Fourier transformation-infrared absorption spectral 20 analysis under the same conditions as those of Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (134), and that the 25 polyhydroxyalkanoate was a copolymer in which an E unit accounted for 9 mol% of the unit.

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 218,000 and a weight average molecular weight Mw of 512,300.

[Example 21]

manner as in Example 19 except that 0.31 g (1.4 mmol) of 2-amino-1-naphthalene sulfonic acid was used instead of 2-aminobenzenesulfonic acid. The resultant polymer was subjected to NMR analysis and Fourier transformation-infrared absorption spectral analysis under the same conditions as those of Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (135), and that the

unit accounted for 8 mol% of the unit.

$$SO_3H$$
 $O H CH_3$
 O

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 165,000 and a weight average molecular weight Mw of 371,300.

[Example 22]

10 Under a nitrogen atmosphere, 0.40 g of the polyhydroxyalkanoate copolymer composed of the unit represented by the chemical formula (108) synthesized in Example 6 (C: 11 mol%, D: 89 mol%) and 0.26 g (1.5 mmol) of 4-aminobenzenesulfonic acid were placed in a 100-ml three-necked flask. 15.0 ml of pyridine was added to the flask, and the mixture was stirred. After that, 0.78 ml (3.0 mmol) of triphenyl phosphite was added. After that, 0.34 g of a polymer was prepared in the same manner as in Example 19. The

Fourier transformation-infrared absorption spectral analysis under the same conditions as those of Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (136), and that the polyhydroxyalkanoate was a copolymer in which an E unit accounted for 10 mol% of the unit.

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 218,000 and a weight average molecular weight Mw of 545,000.

[Example 23]

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0.31 g of a polymer was prepared in the same manner as in Example 22 except that 0.23 g (1.5 mmol) of 2-amino-2-methylpropane sulfonic acid was used instead of 4-aminobenzenesulfonic acid in Example 22.

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The resultant polymer was subjected to NMR analysis and Fourier transformation-infrared absorption spectral analysis under the same conditions as those of Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (137), and that the polyhydroxyalkanoate was a copolymer in which an E unit accounted for 9 mol% of the unit.

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 225,000 and a weight average molecular weight Mw of 540,000.

[Example 24]

0.35 g of a polymer was prepared in the same manner as in Example 22 except that 0.33 g (1.5 mmol) of 1-naphthylamine-8-sulfonic acid was used instead of 4-aminobenzenesulfonic acid in Example 22. The

resultant polymer was subjected to NMR analysis and Fourier transformation-infrared absorption spectral analysis under the same conditions as those of Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (138), and that the polyhydroxyalkanoate was a copolymer in which an E unit accounted for 8 mol% of the unit.

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The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 178,000 and a weight average molecular weight Mw of 445,000.

[Example 25]

0.37 g of a polymer was prepared in the same manner as in Example 22 except that 0.37 g (1.5 mmol) of 2-aminobenzenesulfonic acid phenyl ester was used instead of 4-aminobenzenesulfonic acid in Example 22.

The resultant polymer was subjected to NMR analysis and Fourier transformation-infrared absorption spectral analysis under the same conditions as those of Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (139), and that the polyhydroxyalkanoate was a copolymer in which an E unit accounted for 9 mol% of the unit.

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 210,500 and a weight average molecular weight Mw of 509,400.

[Example 26]

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Under a nitrogen atmosphere, 0.40 g of the polyhydroxyalkanoate copolymer composed of the unit represented by the chemical formula (110) synthesized in Example 7 (C: 10 mol%, D: 90 mol%) and 0.23 g (1.3)

mmol) of 2-aminobenzenesulfonic acid were placed in a 100-ml three-necked flask. 15.0 ml of pyridine was added to the flask, and the mixture was stirred. After that, 0.69 ml (2.7 mmol) of triphenyl phosphite was added. After that, 0.34 g of a polymer was 5 prepared in the same manner as in Example 19. The resultant polymer was subjected to NMR analysis and Fourier transformation-infrared absorption spectral analysis under the same conditions as those of Example 19. As a result, it was confirmed that the 10 resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (140), and that the polyhydroxyalkanoate was a copolymer in which an E unit accounted for 10 mol% of the unit. 15

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular

weight Mn of 208,000 and a weight average molecular weight Mw of 499,200.

[Example 27]

0.29 g of a polymer was prepared in the same
manner as in Example 26 except that 0.17 g (1.3 mmol)
of taurine was used instead of 2-aminobenzenesulfonic
acid in Example 26. The resultant polymer was
subjected to NMR analysis and Fourier transformationinfrared absorption spectral analysis under the same
conditions as those of Example 19. As a result, it
was confirmed that the resultant polymer was a
polyhydroxyalkanoate containing a unit represented by
the following chemical formula (141), and that the
polyhydroxyalkanoate was a copolymer in which an E
unit accounted for 7 mol% of the unit.

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular

weight Mn of 225,000 and a weight average molecular weight Mw of 562,500.

[Example 28]

Under a nitrogen atmosphere, 0.40 g of the polyhydroxyalkanoate copolymer composed of the unit 5 represented by the chemical formula (112) synthesized in Example 8 (C: 9 mol%, D: 91 mol%) and 0.22 g (1.2 mmol) of p-toluidine-2-sulfonic acid were placed in a 100-ml three-necked flask. 15.0 ml of pyridine was added to the flask, and the mixture was stirred. 10 After that, 0.60 ml (2.3 mmol) of triphenyl phosphite was added. After that, 0.32 g of a polymer was prepared in the same manner as in Example 19. The resultant polymer was subjected to NMR analysis and Fourier transformation-infrared absorption spectral 15 analysis under the same conditions as those of Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (142), and that the 20 polyhydroxyalkanoate was a copolymer in which an E unit accounted for 8 mol% of the unit.

$$CH_3$$
 SO_3H
 $N-H$
 $=0$
 $(CH_2)_{7H}$
 O
 H
 CH_3
 F
 (142)

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 215,500 and a weight average molecular weight Mw of 538,800.

[Example 29]

0.34 g of a polymer was prepared in the same
manner as in Example 28 except that 0.26 g (1.2 mmol)
of 2-amino-1-naphthalene sulfonic acid was used
instead of p-toluidine-2-sulfonic acid in Example 28.
The resultant polymer was subjected to NMR analysis
and Fourier transformation-infrared absorption
spectral analysis under the same conditions as those
of Example 19. As a result, it was confirmed that
the resultant polymer was a polyhydroxyalkanoate
containing a unit represented by the following
chemical formula (143), and that the

polyhydroxyalkanoate was a copolymer in which an E unit accounted for 8 mol% of the unit.

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 198,100 and a weight average molecular weight Mw of 486,300.

10 [Example 30]

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Under a nitrogen atmosphere, 0.40 g of the polyhydroxyalkanoate copolymer composed of the unit represented by the chemical formula (114) synthesized in Example 9 (C: 12 mol%, D: 88 mol%) and 0.23 g (1.3 mmol) of 2-aminobenzenesulfonic acid were placed in a 100-ml three-necked flask. 15.0 ml of pyridine was added to the flask, and the mixture was stirred. After that, 0.69 ml (2.6 mmol) of triphenyl phosphite was added. After that, 0.33 g of a polymer was

prepared in the same manner as in Example 19. The resultant polymer was subjected to NMR analysis and Fourier transformation-infrared absorption spectral analysis under the same conditions as those of

5 Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (144), and that the polyhydroxyalkanoate was a copolymer in which an E

10 unit accounted for 11 mol% of the unit.

$$SO_3H$$
 $O H H H O H CH_2)_2$
 F
 $O H H H O H CH_2)_2$
 $O H CH_2)_2$
 $O H H H O H CH_2)_2$

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 55,300 and a weight average molecular weight Mw of 113,400.

[Example 31]

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0.35 g of a polymer was prepared in the same

manner as in Example 30 except that 0.27 g (1.3 mmol) of 4-methoxyaniline-2-sulfonic acid was used instead of 2-aminobenzenesulfonic acid in Example 30. The resultant polymer was subjected to NMR analysis and 5 Fourier transformation-infrared absorption spectral analysis under the same conditions as those of Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (145), and that the polyhydroxyalkanoate was a copolymer in which an E unit accounted for 11 mol% of the unit.

The average molecular weight of the resultant

polymer was measured under the same conditions as
those of Example 19. As a result, the resultant
polymer was found to have a number average molecular

weight Mn of 56,000 and a weight average molecular weight Mw of 117,600.

[Example 32]

- 0.31 g of a polymer was prepared in the same manner as in Example 30 except that 0.30 g (1.3 mmol) 5 of 2-amino-1-naphthalene sulfonic acid was used instead of 2-aminobenzenesulfonic acid in Example 30. The resultant polymer was subjected to NMR analysis and Fourier transformation-infrared absorption spectral analysis under the same conditions as those 10 of Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (146), and that the polyhydroxyalkanoate was a copolymer in which an E 15 unit accounted for 9 mol% of the unit.

The average molecular weight of the resultant

polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 38,500 and a weight average molecular weight Mw of 82,800.

[Example 33]

Under a nitrogen atmosphere, 0.40 g of the polyhydroxyalkanoate copolymer composed of the unit represented by the chemical formula (116) synthesized in Example 10 (C: 12 mol%, D: 88 mol%) and 0.19 g 10 (1.1 mmol) of 2-aminobenzenesulfonic acid were placed in a 100-ml three-necked flask. 15.0 ml of pyridine was added to the flask, and the mixture was stirred. After that, 0.57 ml (2.2 mmol) of triphenyl phosphite was added. After that, 0.33 g of a polymer was 15 prepared in the same manner as in Example 19. The resultant polymer was subjected to NMR analysis and Fourier transformation-infrared absorption spectral analysis under the same conditions as those of Example 19. As a result, it was confirmed that the 20 resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (147), and that the polyhydroxyalkanoate was a copolymer in which an E unit accounted for 10 mol% of the unit. 25

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 52,500 and a weight average molecular weight Mw of 107,600.

[Example 34]

manner as in Example 33 except that 0.19 g (1.1 mmol) of 3-aminobenzene sulfonic acid was used instead of 2-aminobenzenesulfonic acid in Example 33. The resultant polymer was subjected to NMR analysis and Fourier transformation-infrared absorption spectral analysis under the same conditions as those of Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following

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chemical formula (148), and that the polyhydroxyalkanoate was a copolymer in which an E unit accounted for 9 mol% of the unit.

$$SO_3H$$
 $N-H$
 CH_2H
 O
 H
 $CH_2)_2$
 $CH_2(CH_2)_2$
 $CH_2(CH_2)_2$

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 51,800 and a weight average molecular weight Mw of 108,800.

[Example 35]

0.35 g of a polymer was prepared in the same manner as in Example 33 except that 0.19 g (1.1 mmol) of 4-aminobenzene sulfonic acid was used instead of 2-aminobenzene sulfonic acid in Example 33. The resultant polymer was subjected to NMR analysis and Fourier transformation-infrared absorption spectral analysis under the same conditions as those of

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Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (149), and that the

5 polyhydroxyalkanoate was a copolymer in which an E unit accounted for 10 mol% of the unit.

$$F$$

SO₃H

N-H

CH₂H

O H

(CH₂)₂

F

(149)

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 51,500 and a weight average molecular weight Mw of 103,000.

[Example 36]

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0.37 g of a polymer was prepared in the same manner as in Example 33 except that 0.22 g (1.1 mmol) of 4-methoxyaniline-2-sulfonic acid was used instead

of 2-aminobenzenesulfonic acid in Example 33. The resultant polymer was subjected to NMR analysis and Fourier transformation-infrared absorption spectral analysis under the same conditions as those of

5 Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (150), and that the polyhydroxyalkanoate was a copolymer in which an E unit accounted for 10 mol% of the unit.

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 49,800 and a weight average molecular weight Mw of 102,100.

[Example 37]

manner as in Example 33 except that 0.17 g (1.1 mmol) of 2-amino-2-methylpropane sulfonic acid was used instead of 2-aminobenzenesulfonic acid in Example 33. The resultant polymer was subjected to NMR analysis and Fourier transformation-infrared absorption spectral analysis under the same conditions as those of Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (151), and that the polyhydroxyalkanoate was a copolymer in which an E unit accounted for 8 mol% of the unit.

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant

polymer was found to have a number average molecular weight Mn of 53,200 and a weight average molecular weight Mw of 111,700.

[Example 38]

5 ' Under a nitrogen atmosphere, 0.40 g of the polyhydroxyalkanoate copolymer composed of the unit represented by the chemical formula (118) synthesized in Example 11 (C: 11 mol%, D: 89 mol%) and 0.21 g (1.2 mmol) of 2-aminobenzenesulfonic acid were placed in a 100-ml three-necked flask. 15.0 ml of pyridine 10 was added to the flask, and the mixture was stirred. After that, 0.62 ml (2.4 mmol) of triphenyl phosphite was added. After that, 0.33 g of a polymer was prepared in the same manner as in Example 19. The 15 resultant polymer was subjected to NMR analysis and Fourier transformation-infrared absorption spectral analysis under the same conditions as those of Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following 20 chemical formula (152), and that the polyhydroxyalkanoate was a copolymer in which an E unit accounted for 11 mol% of the unit.

$$SO_3H$$
 $OH_{(CH_2)_2}$
 $OH_{$

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 53,500 and a weight average molecular weight Mw of 104,300.

[Example 39]

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0.33 g of a polymer was prepared in the same
manner as in Example 38 except that 0.27 g (1.2 mmol)
of 2-amino-1-naphthalene sulfonic acid was used
instead of 2-aminobenzenesulfonic acid in Example 38.
The resultant polymer was subjected to NMR analysis
and Fourier transformation-infrared absorption
spectral analysis under the same conditions as those
of Example 19. As a result, it was confirmed that
the resultant polymer was a polyhydroxyalkanoate
containing a unit represented by the following

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chemical formula (153), and that the polyhydroxyalkanoate was a copolymer in which an E unit accounted for 10 mol% of the unit.

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 37,600 and a weight average molecular weight Mw of 77,100.

[Example 40]

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Under a nitrogen atmosphere, 0.40 g of the polyhydroxyalkanoate copolymer composed of the unit represented by the chemical formula (120) synthesized in Example 12 (C: 8 mol%, D: 92 mol%) and 0.15 g (0.9 mmol) of 2-aminobenzenesulfonic acid were placed in a 100-ml three-necked flask. 15.0 ml of pyridine was

added to the flask, and the mixture was stirred.

After that, 0.45 ml (1.7 mmol) of triphenyl phosphite was added. After that, 0.34 g of a polymer was prepared in the same manner as in Example 19. The

5 resultant polymer was subjected to NMR analysis and Fourier transformation-infrared absorption spectral analysis under the same conditions as those of Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate

10 containing a unit represented by the following chemical formula (154), and that the polyhydroxyalkanoate was a copolymer in which an E unit accounted for 7 mol% of the unit.

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular

(154)

weight Mn of 54,200 and a weight average molecular weight Mw of 108,400.

[Example 41]

0.35 g of a polymer was prepared in the same
manner as in Example 40 except that 0.22 g (0.9 mmol)
of 2-aminobenzenesulfonic acid phenyl ester was used
instead of 2-aminobenzenesulfonic acid in Example 40.
The resultant polymer was subjected to NMR analysis
and Fourier transformation-infrared absorption
spectral analysis under the same conditions as those
of Example 19. As a result, it was confirmed that
the resultant polymer was a polyhydroxyalkanoate
containing a unit represented by the following
chemical formula (155), and that the
polyhydroxyalkanoate was a copolymer in which an E
unit accounted for 7 mol% of the unit.

The average molecular weight of the resultant

polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 55,000 and a weight average molecular weight Mw of 104,500.

[Example 42]

Under a nitrogen atmosphere, 0.40 g of the polyhydroxyalkanoate copolymer composed of the unit represented by the chemical formula (122) synthesized 10 in Example 13 (C: 11 mol%, D: 89 mol%) and 0.19 g (1.1 mmol) of 2-aminobenzenesulfonic acid were placed in a 100-ml three-necked flask. 15.0 ml of pyridine was added to the flask, and the mixture was stirred. After that, 0.58 ml (2.2 mmol) of triphenyl phosphite 15 was added. After that, 0.33 g of a polymer was prepared in the same manner as in Example 19. The resultant polymer was subjected to NMR analysis and Fourier transformation-infrared absorption spectral analysis under the same conditions as those of 20 Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (156), and that the polyhydroxyalkanoate was a copolymer in which an E 25 unit accounted for 10 mol% of the unit.

[Example 43]

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The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 100,500 and a weight average molecular weight Mw of 221,100.

0.35 g of a polymer was prepared in the same
manner as in Example 42 except that 0.23 g (1.1 mmol)
of 4-methoxyaniline-2-sulfonic acid was used instead
of 2-aminobenzenesulfonic acid in Example 42. The
resultant polymer was subjected to NMR analysis and
Fourier transformation-infrared absorption spectral

analysis under the same conditions as those of Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following

chemical formula (157), and that the polyhydroxyalkanoate was a copolymer in which an E unit accounted for 7 mol% of the unit.

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 110,200 and a weight average molecular weight Mw of 236,900.

[Example 44]

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Under a nitrogen atmosphere, 0.40 g of the polyhydroxyalkanoate copolymer composed of the unit represented by the chemical formula (124) synthesized in Example 14 (C: 11 mol%, D: 89 mol%) and 0.25 g (1.1 mmol) of 2-amino-1-napthalene sulfonic acid were placed in a 100-ml three-necked flask. 15.0 ml of

pyridine was added to the flask, and the mixture was stirred. After that, 0.58 ml (2.2 mmol) of triphenyl phosphite was added. After that, 0.34 g of a polymer was prepared in the same manner as in Example 19.

5 The resultant polymer was subjected to NMR analysis and Fourier transformation-infrared absorption spectral analysis under the same conditions as those of Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (158), and that the polyhydroxyalkanoate was a copolymer in which an E

unit accounted for 11 mol% of the unit.

The average molecular weight of the resultant polymer was measured under the same conditions as

those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 87,500 and a weight average molecular weight Mw of 192,500.

5 [Example 45]

manner as in Example 44 except that 0.17 g (1.1 mmol) of 2-amino-2-methylpropane sulfonic acid was used instead of 2-aminobenzenesulfonic acid in Example 44.

The resultant polymer was subjected to NMR analysis and Fourier transformation-infrared absorption spectral analysis under the same conditions as those of Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (159), and that the polyhydroxyalkanoate was a copolymer in which an E unit accounted for 7 mol% of the unit.

[Example 46]

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 99,800 and a weight average molecular weight Mw of 214,600.

Under a nitrogen atmosphere, 0.40 g of the

10 polyhydroxyalkanoate copolymer composed of the unit
represented by the chemical formula (128) synthesized
in Example 16 (C: 8 mol%, D: 92 mol%) and 0.15 g (0.9
mmol) of 2-aminobenzenesulfonic acid were placed in a
100-ml three-necked flask. 15.0 ml of pyridine was
15 added to the flask, and the mixture was stirred.
After that, 0.46 ml (1.8 mmol) of triphenyl phosphite
was added. After that, 0.33 g of a polymer was

prepared in the same manner as in Example 19. The resultant polymer was subjected to NMR analysis and Fourier transformation-infrared absorption spectral analysis under the same conditions as those of

5 Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (160), and that the polyhydroxyalkanoate was a copolymer in which an E

10 unit accounted for 8 mol% of the unit.

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 100,100 and a weight average molecular weight Mw of 225,200.

[Example 47]

manner as in Example 46 except that 0.22 g (0.9 mmol) of 4-aminobenezenesulfonic acid phenyl ester was used instead of 2-aminobenzenesulfonic acid in Example 46. The resultant polymer was subjected to NMR analysis and Fourier transformation-infrared absorption spectral analysis under the same conditions as those of Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (161), and that the polyhydroxyalkanoate was a copolymer in which an E unit accounted for 7 mol% of the unit.

$$\begin{array}{c|c}
\hline
O = S = O \\
\hline
N - H \\
\hline
O \\
O + H \\
\hline
H (CH_2)_2
\end{array}$$

$$\begin{array}{c|c}
O & H & H \\
\hline
H (CH_2)_2
\end{array}$$

$$\begin{array}{c|c}
F \\
\end{array}$$
(161)

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 110,500 and a weight average molecular weight Mw of 237,600.

[Example 48]

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Under a nitrogen atmosphere, 0.40 g of the

10 polyhydroxyalkanoate copolymer composed of the unit
represented by the chemical formula (130) synthesized
in Example 17 (C: 10 mol%, D: 90 mol%) and 0.26 g

(1.2 mmol) of 2-amino-1-naphthalene sulfonic acid

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were placed in a 100-ml three-necked flask. 15.0 ml of pyridine was added to the flask, and the mixture was stirred. After that, 0.60 ml (2.3 mmol) of triphenyl phosphite was added. After that, 0.36 g of a polymer was prepared in the same manner as in Example 19. The resultant polymer was subjected to NMR analysis and Fourier transformation-infrared absorption spectral analysis under the same conditions as those of Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (162), and that the polyhydroxyalkanoate was a copolymer in which an E unit accounted for 9 mol% of the unit.

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant

polymer was found to have a number average molecular weight Mn of 30,500 and a weight average molecular weight Mw of 65,600.

[Example 49]

Under a nitrogen atmosphere, 0.40 g of the 5 polyhydroxyalkanoate copolymer composed of the unit represented by the chemical formula (132) synthesized in Example 18 (C: 9 mol%, D: 91 mol%) and 0.16 g (1.0 mmol) of 2-amino-2-methylpropane sulfonic acid were placed in a 100-ml three-necked flask. 15.0 ml of 10 pyridine was added to the flask, and the mixture was stirred. After that, 0.52 ml (2.0 mmol) of triphenyl phosphite was added. After that, 0.31 g of a polymer was prepared in the same manner as in Example 19. The resultant polymer was subjected to NMR analysis 1.5 and Fourier transformation-infrared absorption spectral analysis under the same conditions as those of Example 19. As a result, it was confirmed that the resultant polymer was a polyhydroxyalkanoate 20 containing a unit represented by the following chemical formula (163), and that the polyhydroxyalkanoate was a copolymer in which an E unit accounted for 8 mol% of the unit.

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 19. As a result, the resultant polymer was found to have a number average molecular weight Mn of 32,500 and a weight average molecular weight Mw of 71,500.

[Example 50]

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0.30 g of the polyhydroxyalkanoate copolymer

composed of the unit represented by the chemical formula (133) synthesized in Example 19 was added to a round-bottomed flask. Then, 21.0 ml of chloroform and 7.0 ml of methanol were added to dissolve the polymer, and the solution was cooled to 0°C. 0.93 ml of a 2-mol/L trimethylsilyldiazomethane-hexane solution (manufactured by Aldrich) was added to the solution, and the whole was stirred for 4 hours.

After the completion of the reaction, the solvent was

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distilled off by using an evaporator, and then the polymer was collected. Furthermore, 21.0 ml of chloroform and 7.0 ml of methanol were added to dissolve the polymer again. Then, the solvent was distilled off by using an evaporator. This operation was repeated 3 times. The collected polymer was dried under reduced pressure to prepare 0.30 g of a polymer. The structure of the resultant polymer was determined through analysis according to 1H-NMR (FT-NMR: Bruker DPX 400; resonance frequency: 400 MHz; measured nuclear species: ¹H; solvent used: DMSO-d₆; measurement temperature: room temperature). $^{1}\text{H-NMR}$ confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (164) as a monomer unit because a peak derived from methyl sulfonate was observed at 3 to 4 ppm.

SO₃CH₃

$$O = H$$

$$O$$

It was also confirmed that a G unit accounted for 10 mol% of the unit of the polyhydroxyalkanoate represented by the chemical formula (164). In

addition, there was no peak observed resulted from sulfonic acid in acid value titration using Potentiometric Titrator AT510 (product of Kyoto Electronics Manufacturing Co., Ltd.) and it was also made evident from this that sulfonic acid was 5 converted to methyl sulfonate. The resultant polymer was evaluated for average molecular weight by means of gel permeation chromatography (GPC; Tosoh Corporation HLC-8120, column; Polymer Laboratories PLgel 5 μ MIXED-C, solvent; DMF/LiBr 0.1% (w/v), in 10 terms of polystyrene). As a result, the resultant polymer was found to have a number average molecular weight Mn of 228,000 and a weight average molecular weight Mw of 513,000.

15 [Example 51]

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manner as in Example 50 except that the polyhydroxyalkanoate represented by the chemical formula (134) synthesized in Example 20 was used instead of the polyhydroxyalkanoate represented by the chemical formula (133) synthesized in Example 50, and 0.83 ml of a 2-mol/L trimethylsilyldiazomethane-hexane solution (manufactured by Aldrich) was used. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 50. The analysis confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented

by the following chemical formula (165), and that the polyhydroxyalkanoate was a copolymer in which a G unit accounted for 9 mol% of the unit.

In addition, acid value titration in the same

manner as in Example 50 revealed that the sulfonic
acid was transformed into methyl sulfonate because no
peak derived from the sulfonic acid was observed.

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 50. As a result, the resultant polymer was found to have a number average molecular weight Mn of 209,000 and a weight average molecular weight Mw of 480,700.

15 [Example 52]

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0.30 g of a polymer was prepared in the same manner as in Example 50 except that the polyhydroxyalkanoate represented by the chemical formula (135) synthesized in Example 21 was used instead of the polyhydroxyalkanoate represented by

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the chemical formula (133) synthesized in Example 50, and 0.71 ml of a 2-mol/L trimethylsilyldiazomethane-hexane solution (manufactured by Aldrich) was used. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 50. The analysis confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (166), and that the polyhydroxyalkanoate was a copolymer in which a G unit accounted for 8 mol% of the unit.

In addition, acid value titration in the same manner as in Example 50 revealed that the sulfonic acid was transformed into methyl sulfonate because no peak derived from the sulfonic acid was observed.

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 50. As a result, the resultant polymer was found to have a number average molecular weight Mn of 162,500 and a weight average molecular

weight Mw of 373,800. [Example 53]

0.29 g of a polymer was prepared in the same manner as in Example 50 except that the polyhydroxyalkanoate represented by the chemical 5 formula (137) synthesized in Example 23 was used instead of the polyhydroxyalkanoate represented by the chemical formula (133) synthesized in Example 50, and 0.83 ml of a 2-mol/L trimethylsilyldiazomethanehexane solution (manufactured by Aldrich) was used. 10 The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 50. The analysis confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (167), and that the 15 polyhydroxyalkanoate was a copolymer in which a G unit accounted for 9 mol% of the unit.

In addition, acid value titration in the same manner as in Example 50 revealed that the sulfonic acid was transformed into methyl sulfonate because no peak derived from the sulfonic acid was observed.

$$G \qquad H_{3}C \xrightarrow{CH_{2}} CH_{3}$$

$$H_{3}C \xrightarrow{CH_{2}} CH_{3}$$

$$H_{3}C \xrightarrow{CH_{2}} CH_{3}$$

$$G \qquad H \qquad H$$

$$H \qquad (167)$$

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 50. As a result, the resultant polymer was found to have a number average molecular weight Mn of 228,500 and a weight average molecular weight Mw of 548,400.

[Example 54]

0.30 g of a polymer was prepared in the same manner as in Example 50 except that the 10 polyhydroxyalkanoate represented by the chemical formula (142) synthesized in Example 28 was used instead of the polyhydroxyalkanoate represented by the chemical formula (133) synthesized in Example 50, 15 and 0.71 ml of a 2-mol/L trimethylsilyldiazomethanehexane solution (manufactured by Aldrich) was used. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 50. The analysis confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented 20 by the following chemical formula (168), and that the

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polyhydroxyalkanoate was a copolymer in which a G unit accounted for 8 mol% of the unit.

In addition, acid value titration in the same manner as in Example 50 revealed that the sulfonic acid was transformed into methyl sulfonate because no peak derived from the sulfonic acid was observed.

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 50. As a result, the resultant polymer was found to have a number average molecular weight Mn of 218,000 and a weight average molecular weight Mw of 555,900.

[Example 55]

0.29 g of a polymer was prepared in the same manner as in Example 50 except that the polyhydroxyalkanoate represented by the chemical formula (144) synthesized in Example 30 was used instead of the polyhydroxyalkanoate represented by

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the chemical formula (133) synthesized in Example 50, and 0.83 ml of a 2-mol/L trimethylsilyldiazomethane-hexane solution (manufactured by Aldrich) was used. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 50. The analysis confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (169), and that the polyhydroxyalkanoate was a copolymer in which a G unit accounted for 11 mol% of the unit.

In addition, acid value titration in the same manner as in Example 50 revealed that the sulfonic acid was transformed into methyl sulfonate because no peak derived from the sulfonic acid was observed.

SO₃CH₃

$$O H H O H (CH2)2$$

$$G H H O H (CH2)2$$

$$O H G H (CH2)2$$

$$O H (CH2)2$$

$$O H G H (CH2)2$$

$$O H (CH2)3$$

$$O H (CH2)4$$

$$O H (CH2)2$$

$$O H (CH2)3$$

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 50. As a result, the resultant polymer was found to have a number average molecular

weight Mn of 54,500 and a weight average molecular weight Mw of 114,500.

[Example 56]

0.29 g of a polymer was prepared in the same manner as in Example 50 except that the 5 polyhydroxyalkanoate represented by the chemical formula (149) synthesized in Example 35 was used instead of the polyhydroxyalkanoate represented by the chemical formula (133) synthesized in Example 50, and 0.76 ml of a 2-mol/L trimethylsilyldiazomethane-10 hexane solution (manufactured by Aldrich) was used. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 50. The analysis confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented 15 by the following chemical formula (170), and that the polyhydroxyalkanoate was a copolymer in which a G unit accounted for 10 mol% of the unit.

In addition, acid value titration in the same
20 manner as in Example 50 revealed that the sulfonic
acid was transformed into methyl sulfonate because no
peak derived from the sulfonic acid was observed.

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 50. As a result, the resultant polymer was found to have a number average molecular weight Mn of 51,000 and a weight average molecular weight Mw of 104,600.

[Example 57]

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0.30 g of a polymer was prepared in the same
manner as in Example 50 except that the
polyhydroxyalkanoate represented by the chemical
formula (154) synthesized in Example 40 was used
instead of the polyhydroxyalkanoate represented by
the chemical formula (133) synthesized in Example 50,
and 0.54 ml of a 2-mol/L trimethylsilyldiazomethanehexane solution (manufactured by Aldrich) was used.
The resultant polymer was subjected to NMR analysis

under the same conditions as those of Example 50. The analysis confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (171), and that the polyhydroxyalkanoate was a copolymer in which a G unit accounted for 7 mol% of the unit.

In addition, acid value titration in the same manner as in Example 50 revealed that the sulfonic acid was transformed into methyl sulfonate because no peak derived from the sulfonic acid was observed.

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 50. As a result, the resultant polymer was found to have a number average molecular weight Mn of 52,500 and a weight average molecular weight Mw of 110,300.

[Example 58]

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manner as in Example 50 except that the polyhydroxyalkanoate represented by the chemical formula (156) synthesized in Example 42 was used instead of the polyhydroxyalkanoate represented by the chemical formula (133) synthesized in Example 50, and 0.71 ml of a 2-mol/L trimethylsilyldiazomethane-hexane solution (manufactured by Aldrich) was used. The resultant polymer was subjected to NMR analysis under the same conditions as those of Example 50. The analysis confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (172), and that the polyhydroxyalkanoate was a copolymer in which a G unit accounted for 10 mol% of the unit.

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In addition, oxidation titration in the same manner as in Example 50 revealed that the sulfonic acid was transformed into methyl sulfonate because no peak derived from the sulfonic acid was observed.

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 50. As a result, the resultant polymer was found to have a number average molecular weight Mn of 101,000 and a weight average molecular weight Mw of 227,300.

[Example 59]

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0.29 g of a polymer was prepared in the same
manner as in Example 50 except that the
polyhydroxyalkanoate represented by the chemical
formula (156) synthesized in Example 44 was used
instead of the polyhydroxyalkanoate represented by
the chemical formula (133) synthesized in Example 50,
and 0.75 ml of a 2-mol/L trimethylsilyldiazomethanehexane solution (manufactured by Aldrich) was used.
The resultant polymer was subjected to NMR analysis
under the same conditions as those of Example 50.

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The analysis confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (173), and that the polyhydroxyalkanoate was a copolymer in which a G unit accounted for 11 mol% of the unit.

In addition, acid value titration in the same manner as in Example 50 revealed that the sulfonic acid was transformed into methyl sulfonate because no peak derived from the sulfonic acid was observed.

The average molecular weight of the resultant polymer was measured under—the same conditions as those of Example 50. As a result, the resultant polymer was found to have a number average molecular weight Mn of 86,500 and a weight average molecular weight Mw of 186,000.

[Example 60]

0.28 g of a polymer was prepared in the same manner as in Example 50 except that the polyhydroxyalkanoate represented by the chemical formula (162) synthesized in Example 48 was used 5 instead of the polyhydroxyalkanoate represented by the chemical formula (133) synthesized in Example 50, and 0.71 ml of a 2-mol/L trimethylsilyldiazomethanehexane solution (manufactured by Aldrich) was used. The resultant polymer was subjected to NMR analysis 10 under the same conditions as those of Example 50. The analysis confirmed that the resultant polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (174), and that the polyhydroxyalkanoate was a copolymer in which a G 15 unit accounted for 9 mol% of the unit.

In addition, acid value titration in the same manner as in Example 50 revealed that the sulfonic acid was transformed into methyl sulfonate because no peak derived from the sulfonic acid was observed.

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 50. As a result, the resultant polymer was found to have a number average molecular weight Mn of 31,000 and a weight average molecular weight Mw of 68,200.

[Example 61]

2.00 g of the polyhydroxyalkanoate composed of
the unit represented by the chemical formula (101)
synthesized in Example 1 was placed in a roundbottomed flask, and 100 ml of THF was added to
dissolve this. The flask was placed under a nitrogen

atmosphere, and the solution was stirred at -78°C.

Next, 11.62 ml of a solution of 2 M of lithium diisopropylamide in THF was gradually added to the flask, and the whole was stirred at -78°C for 30 minutes. Next, 10.19 g of 2-acrylamide-2-

methylpropanesulfonic acid methyl ester were added to the flask, and the whole was stirred at room temperature for 30 minutes. After the completion of the reaction, the reaction solution was poured into 400 ml of an aqueous solution of ammonium chloride, 5 and 200 ml of dichloromethane were added to extract the organic layer. The extracted organic layer was washed with 100 ml of water 3 times. After the organic layer had been collected, the solvent was distilled off to collect a crude polymer. Next, the 10 polymer was dissolved into 12 ml of THF, then dissolved into THF, and reprecipitated in methanol in an amount 50 times that of THF necessary for the dissolution. The precipitate was collected and dried under reduced pressure to prepare 1.63 g of a polymer. 15 The structure of the resultant polymer was determined through analysis according to 1H-NMR (FT-NMR: Bruker DPX 400; resonance frequency: 400 MHz; measured nuclear species: 1H; solvent used: DMSO-d6; measurement temperature: room temperature). 20 analysis confirmed that the polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (175) as a monomer unit. The analysis also confirmed that an E unit accounted for 9 mol% of the monomer unit and an F 25 unit accounted for 91 mol% thereof.

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The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 50. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 286,500 and a weight average molecular weight Mw of 572,500. [Example 62]

2.00 g of the polyhydroxyalkanoate composed of the unit represented by the chemical formula (102) 10 synthesized in Example 2 was placed in a roundbottomed flask, and 100 ml of THF was added to dissolve this. The flask was placed under a nitrogen atmosphere, and the solution was stirred at -78°C. Next, 5.68 ml of a solution of 2 M of lithium 15 diisopropylamide in THF was gradually added to the flask, and the whole was stirred at -78°C for 30 minutes. Next, 4.98 g of 2-acrylamide-2methylpropanesulfonic acid methyl ester was added to the flask, and the whole was stirred at room

temperature for 30 minutes. After the completion of the reaction, the reaction solution was poured into 400 ml of an aqueous solution of ammonium chloride, and 200 ml of dichloromethane was added to extract 5 the organic layer. The extracted organic layer was washed with 100 ml of water 3 times. After the organic layer had been collected, the solvent was distilled off to collect a crude polymer. Next, the polymer was dissolved into 12 ml of THF, then dissolved into THF, and reprecipitated in methanol in 10 an amount 50 times that of THF necessary for the dissolution. The precipitate was collected and dried under reduced pressure to prepare 1.22 g of a polymer. The structure of the resultant polymer was determined through analysis according to ¹H-NMR (FT-NMR: Bruker 15 DPX 400; resonance frequency: 400 MHz; measured nuclear species: ¹H; solvent used: DMSO-d₆; measurement temperature: room temperature). analysis confirmed that the polymer was a polyhydroxyalkanoate containing a unit represented by 20 the following chemical formula (176) as a monomer unit. The analysis also confirmed that an E unit accounted for 8 mol% of the monomer unit and an F unit accounted for 92 mol% thereof.

The average molecular weight of the resultant polymer was measured under the same conditions as those of Example 50. As a result, the resultant polyhydroxyalkanoate was found to have a number average molecular weight Mn of 56,500 and a weight average molecular weight Mw of 112,300.

INDUSTRIAL APPLICABILITY

According to the present invention, there is provided: a novel polyhydroxyalkanoate containing, in a molecule, a carboxyl group as a reaction active group at a side chain thereof; a novel polyhydroxyalkanoate containing, in a molecule, a unit having an amide group and a sulfonic group; and a method of producing such a polyhydroxyalkanoate. A novel polyhydroxyalkanoate having a carboxyl group

can find applications in functional materials because it can introduce a functional group which provides functionalities by using its reaction active group. Furthermore, a polyhydroxyalkanoate containing, in a molecule, a unit having a carboxyl group, an amide group, or a sulfonic group is expected to find use in applications including medical soft members because it is excellent in melt processability, and is excellent in biocompatibility by virtue of its hydrophilicity.

This application claims priority from Japanese Patent Application Nos. 2004-174783 filed June 11, 2004 and 2005-168916 filed June 8, 2005, which are hereby incorporated by reference herein.